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United States Patent [19]

Spindt et al.

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[45] Date of Patent: **Oct. 27, 1998**

[54] **USE OF EARLY FORMED LIFT-OFF LAYER IN FABRICATING GATED ELECTRON-EMITTING DEVICES**

[75] Inventors: **Christopher J. Spindt**, Menlo Park; **John M. Macaulay**, Palo Alto; **Robert M. Duboc, Jr.**, Menlo Park, all of Calif.; **Peter C. Searson**, Baltimore, Md.

[73] Assignee: **Candescent Technologies Corporation**, San Jose, Calif.

[21] Appl. No.: **568,885**

[22] Filed: **Dec. 7, 1995**

Related U.S. Application Data

[63] Continuation of Ser. No. 269,229, Jun. 29, 1994, Pat. No. 5,564,959, which is a continuation-in-part of Ser. No. 118,490, Sep. 8, 1993, Pat. No. 5,462,467, and a continuation-in-part of Ser. No. 158,102, Nov. 24, 1993, Pat. No. 5,559,389.

[51] **Int. Cl.⁶** **H01J 9/02**

[52] **U.S. Cl.** **445/24; 445/50**

[58] **Field of Search** **445/24, 50; 313/309, 313/336**

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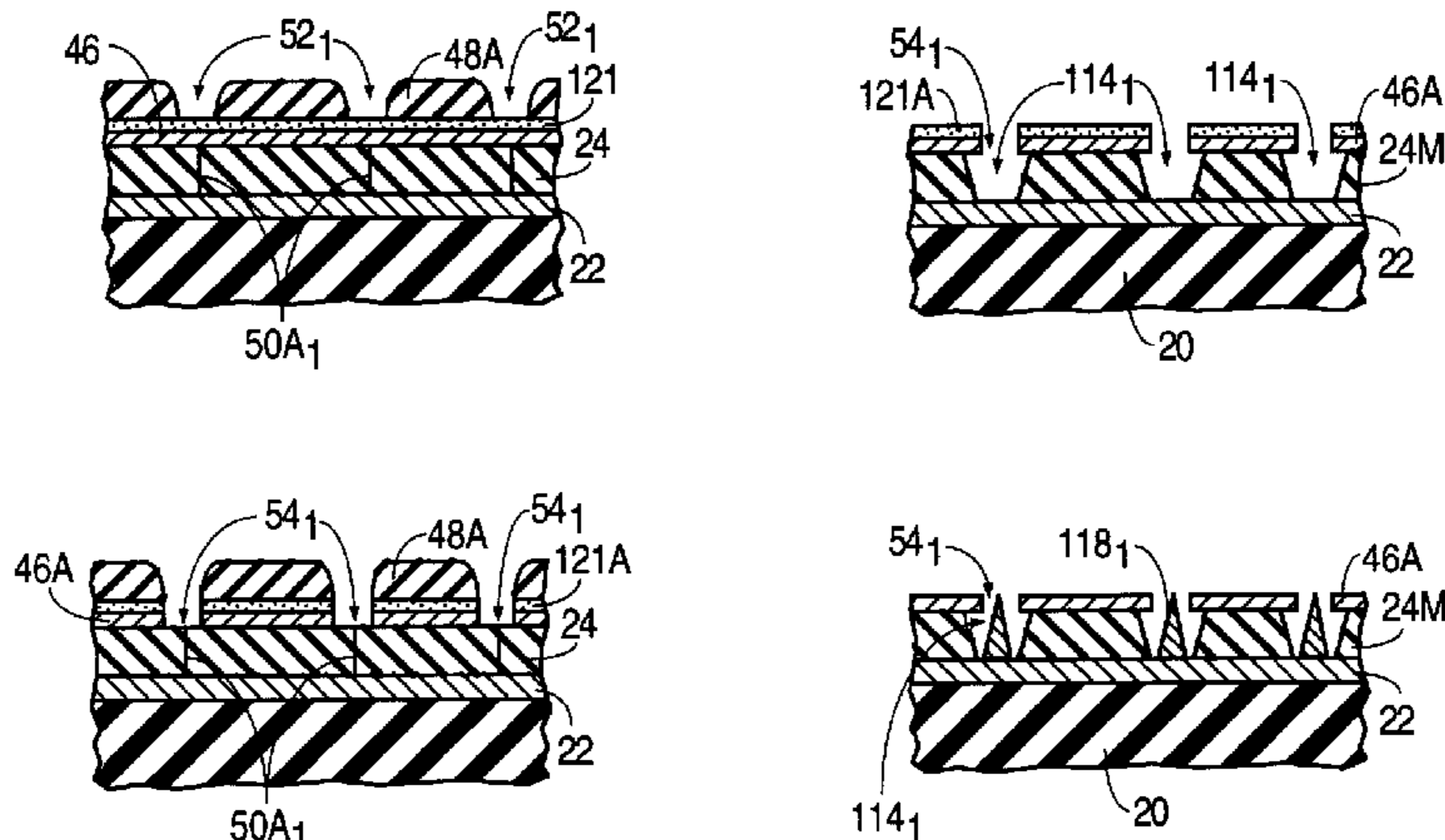
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Primary Examiner—Kenneth J. Ramsey
Attorney, Agent, or Firm—Skjerven, Morrill, MacPherson, Franklin & Friel LLP; Ronald J. Meetin

[57] ABSTRACT

Gated electron emitters are fabricated by processes in which charged particles are passed through a track layer (24, 48, or 144) to form charged-particle tracks (26₁, 50₁, or 146₁). The track layer is etched along the tracks to create open spaces (28₁, 52₁, or 148₁). Electron-emissive elements (30 or 142D) can then be formed at locations respectively centered on the open spaces after which a patterned gate layer (34B, 40B, or 158C) is provided. Alternatively, the open spaces in the track layer can be employed to etch corresponding apertures (54₁) through an underlying non-insulating layer (46) which typically serves as the gate layer. An etch is performed through the apertures to form dielectric open spaces (56₁, 96₁, or 114₁) in an insulating layer (24) that lies below the non-insulating layer. Electron-emissive elements (30B, 30/88D₁, 98/102₁, or 118₁) can subsequently be provided, typically in the dielectric open spaces.

18 Claims, 19 Drawing Sheets



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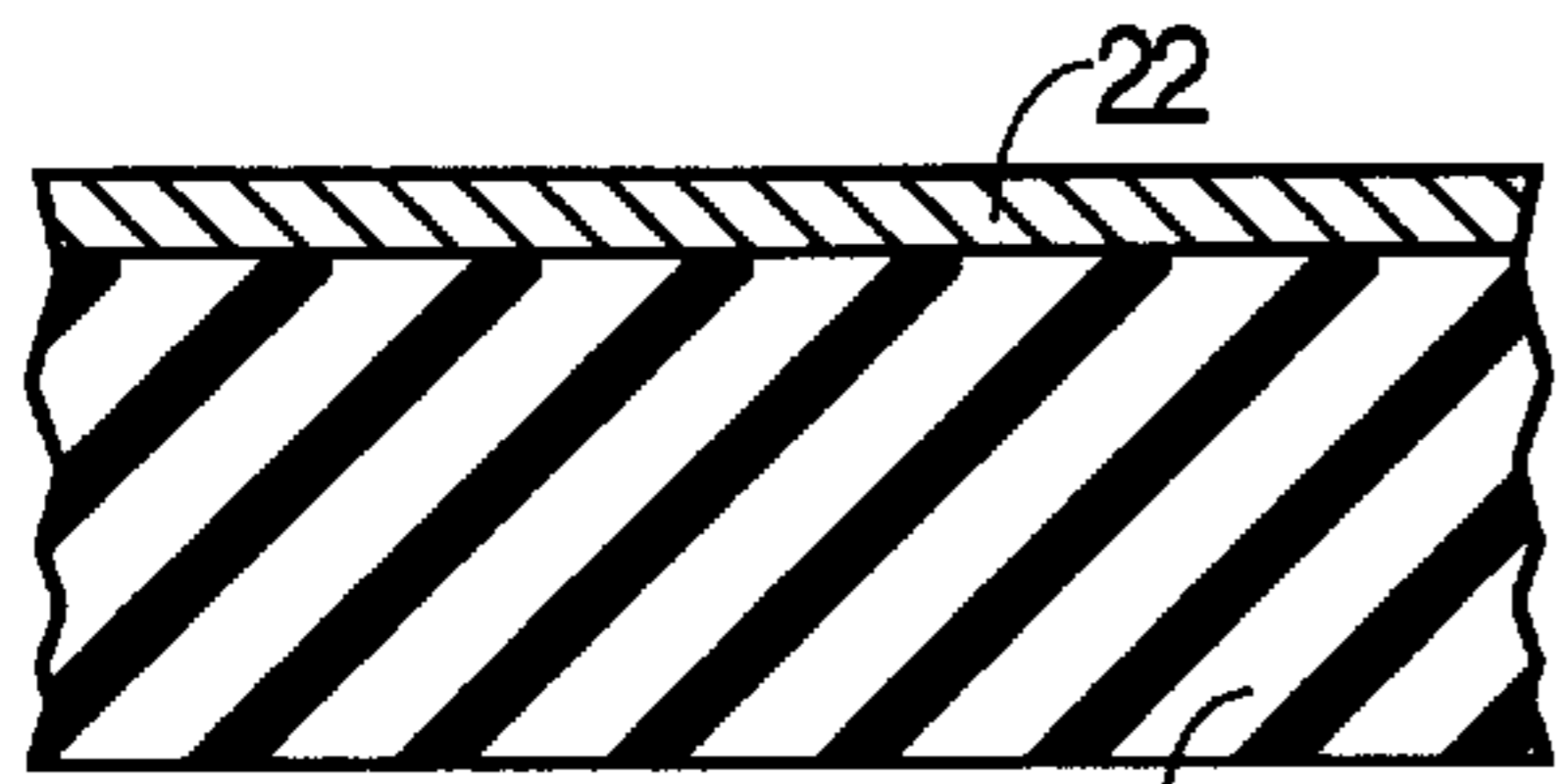


FIG. 1a

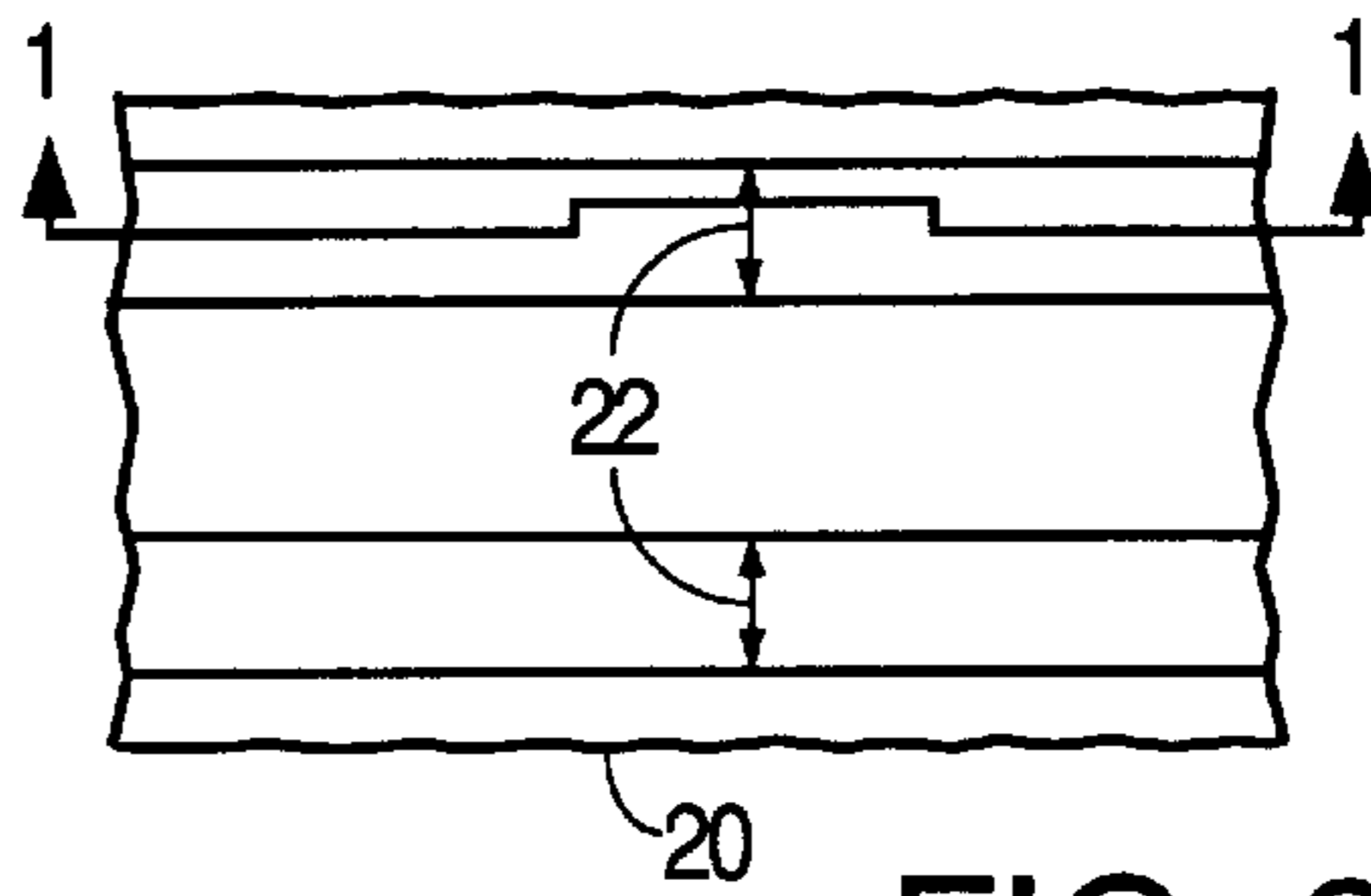


FIG. 2a

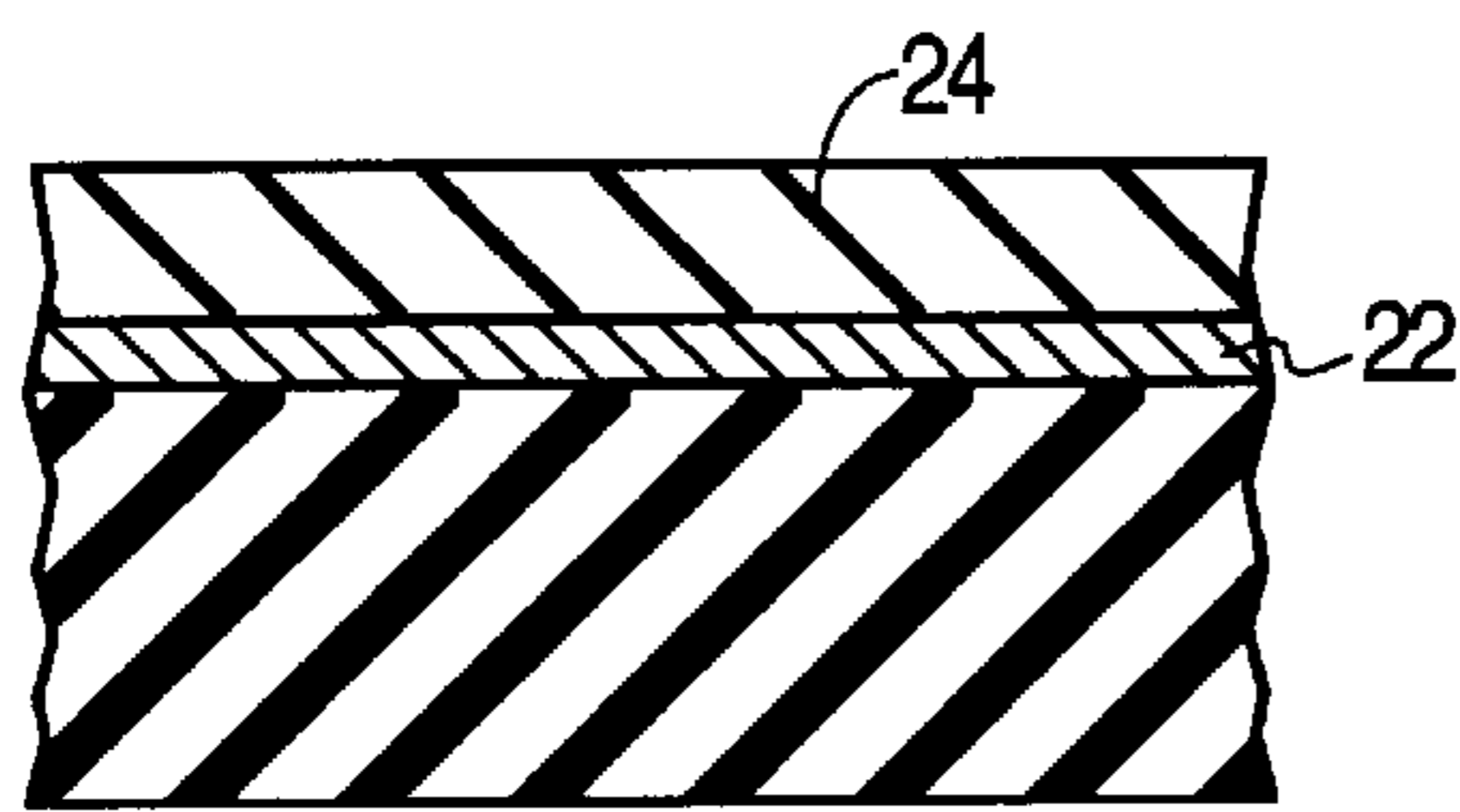


FIG. 1b

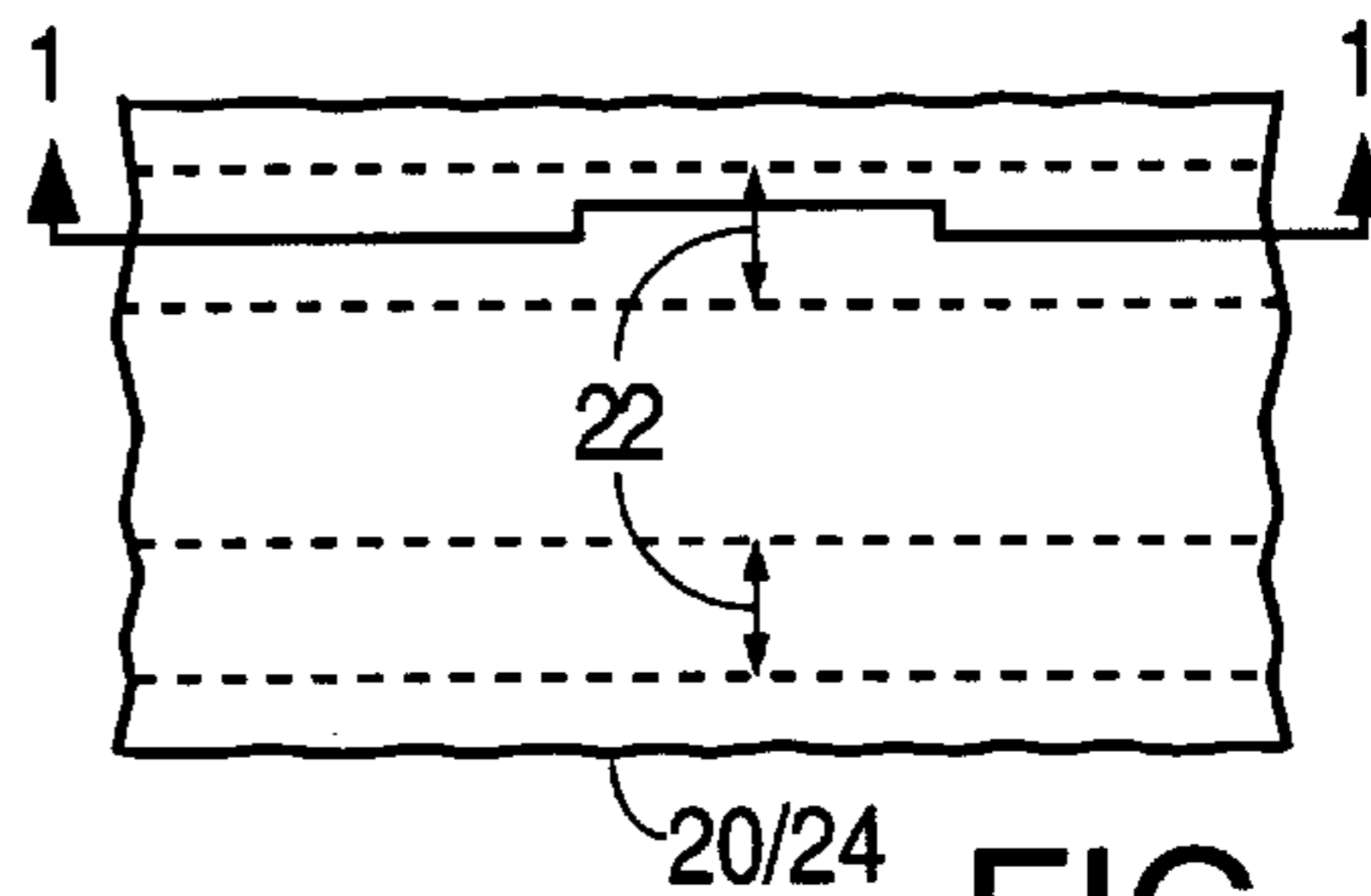


FIG. 2b

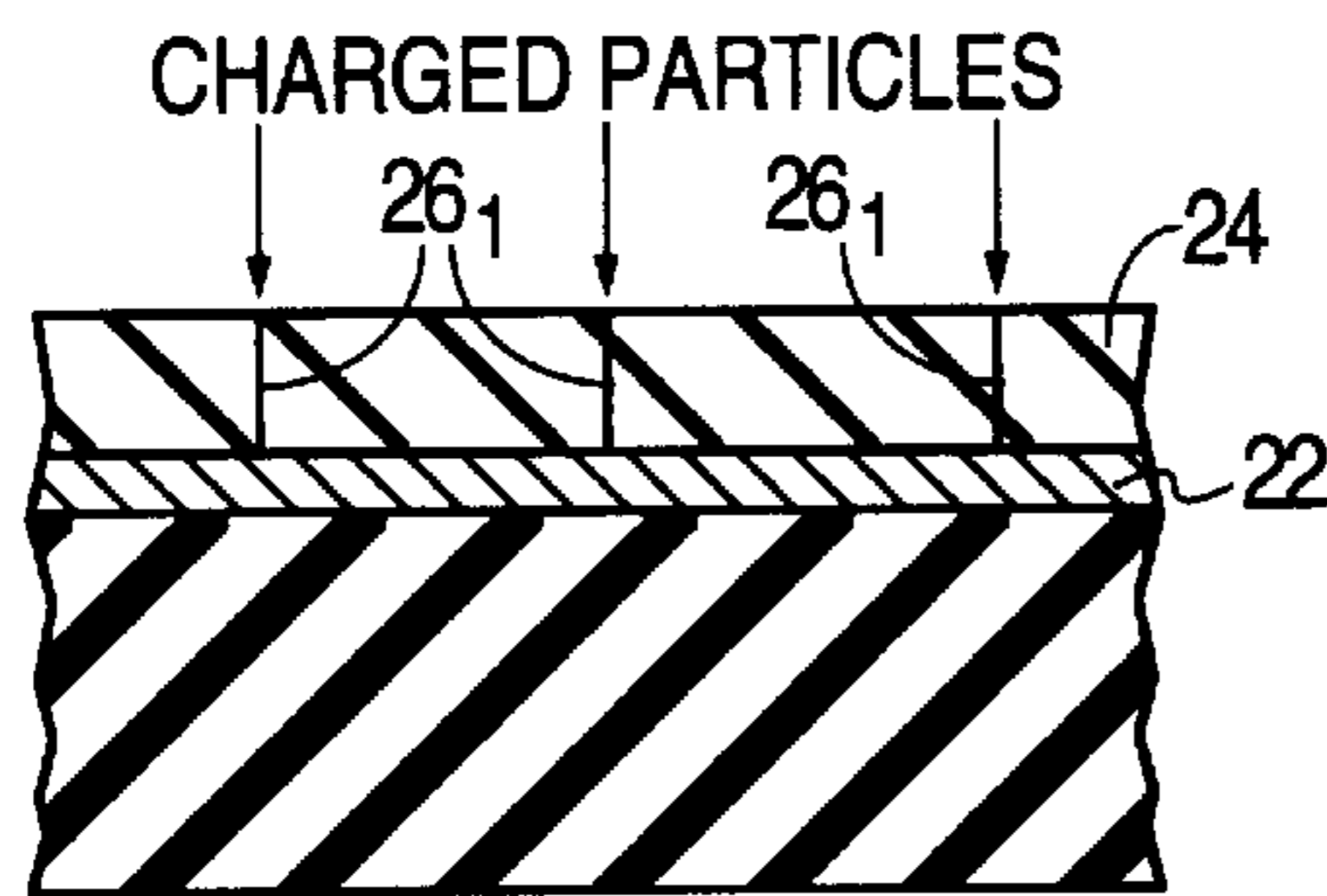


FIG. 1c

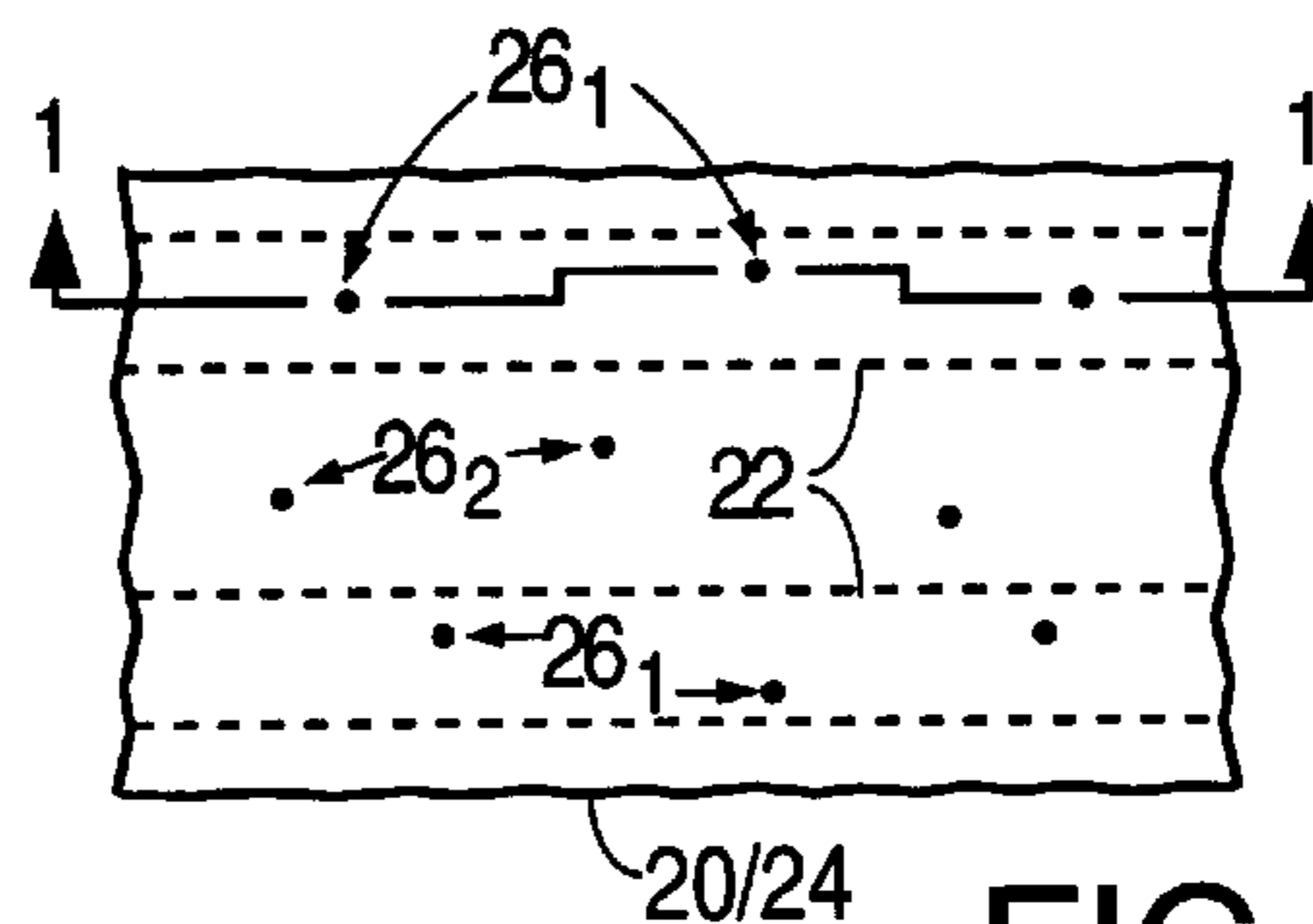


FIG. 2c

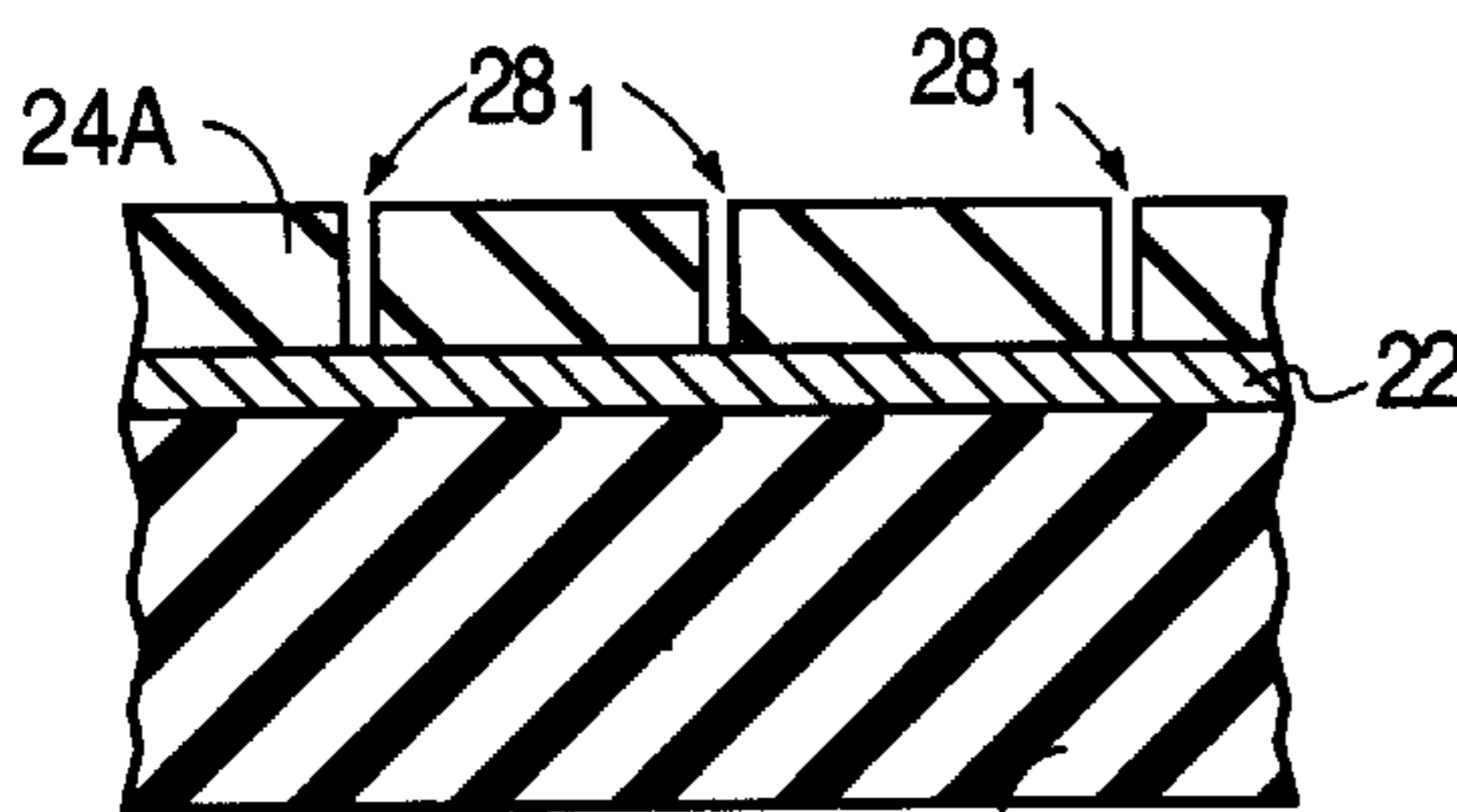


FIG. 1d

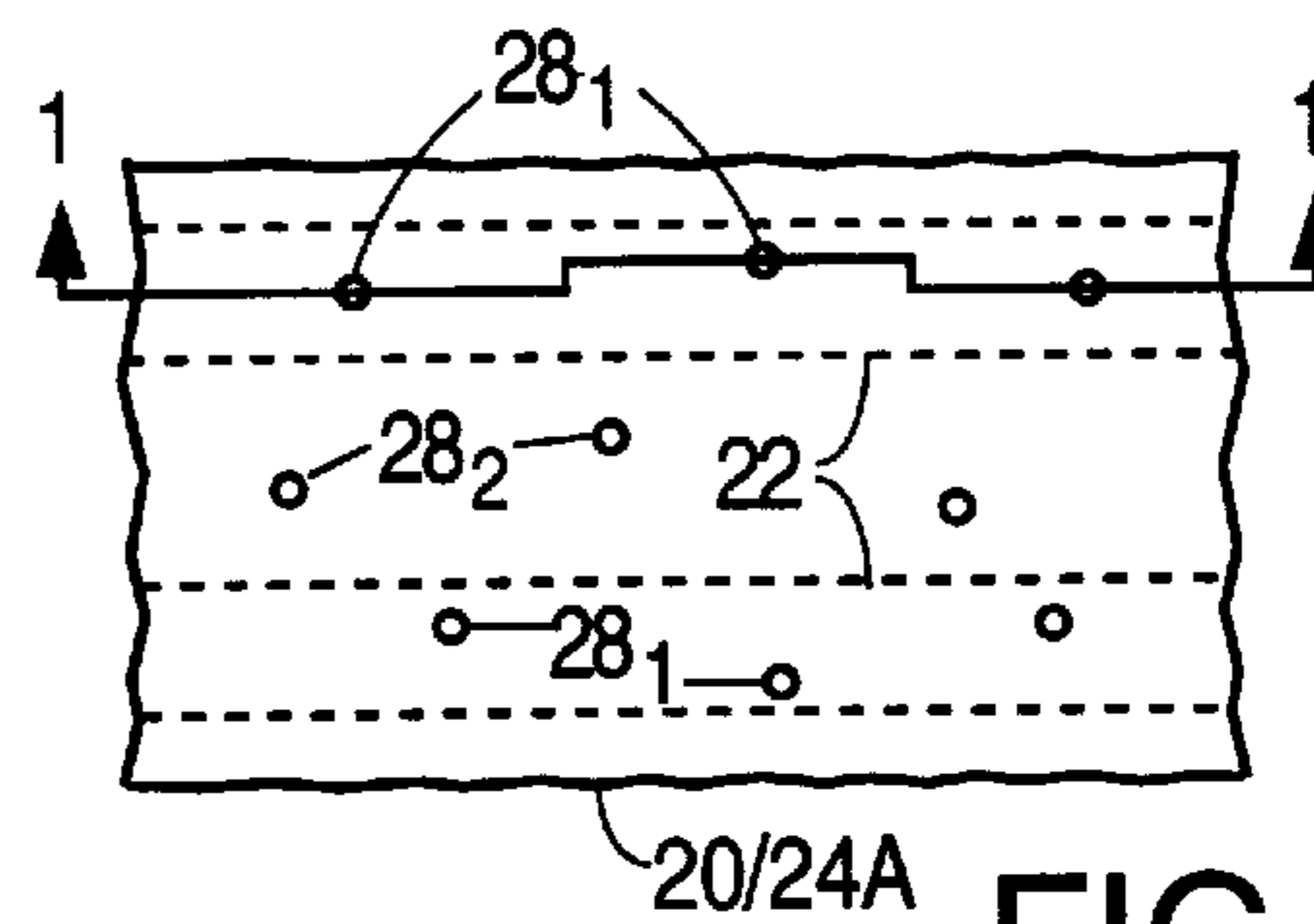


FIG. 2d

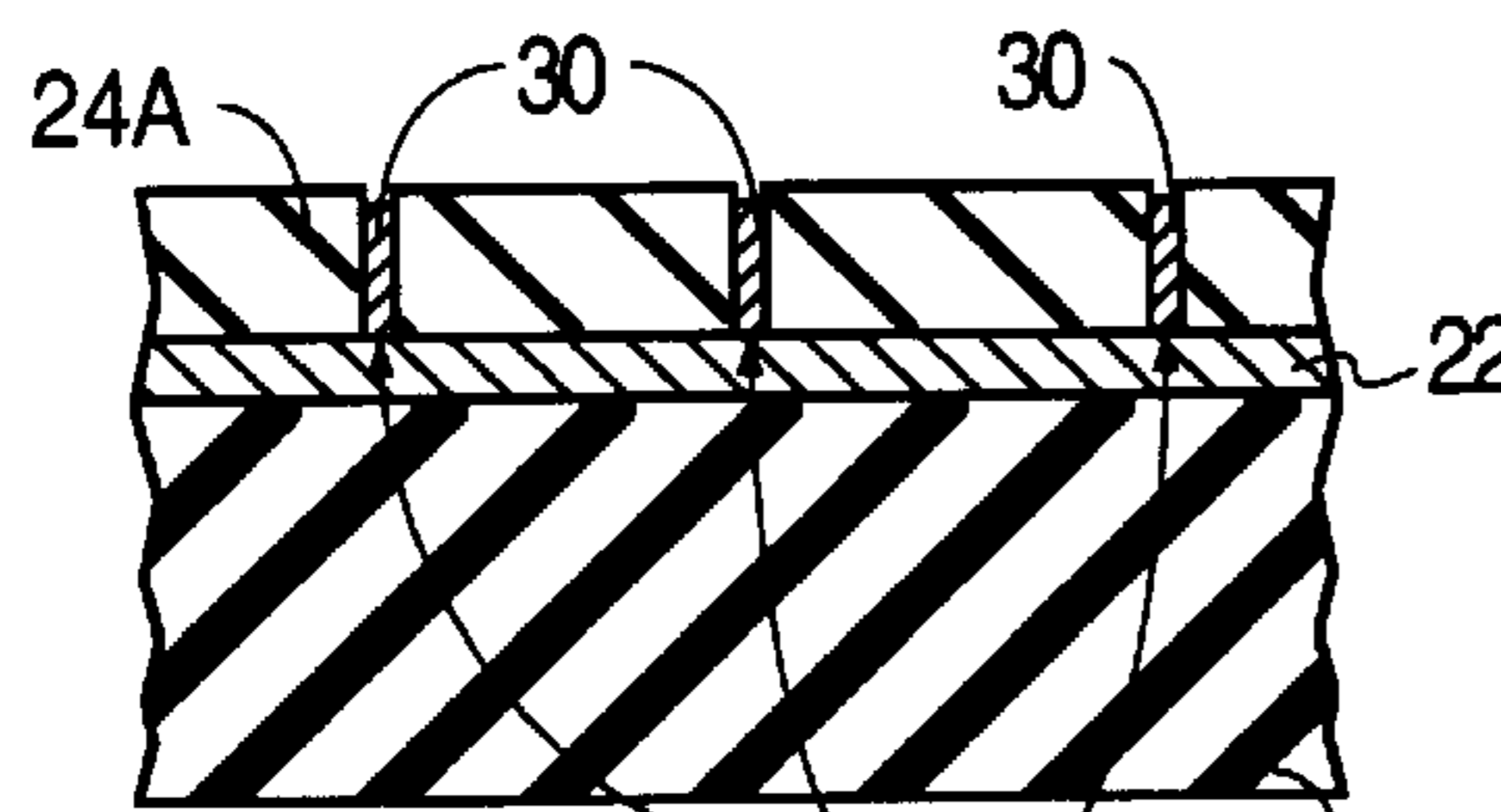


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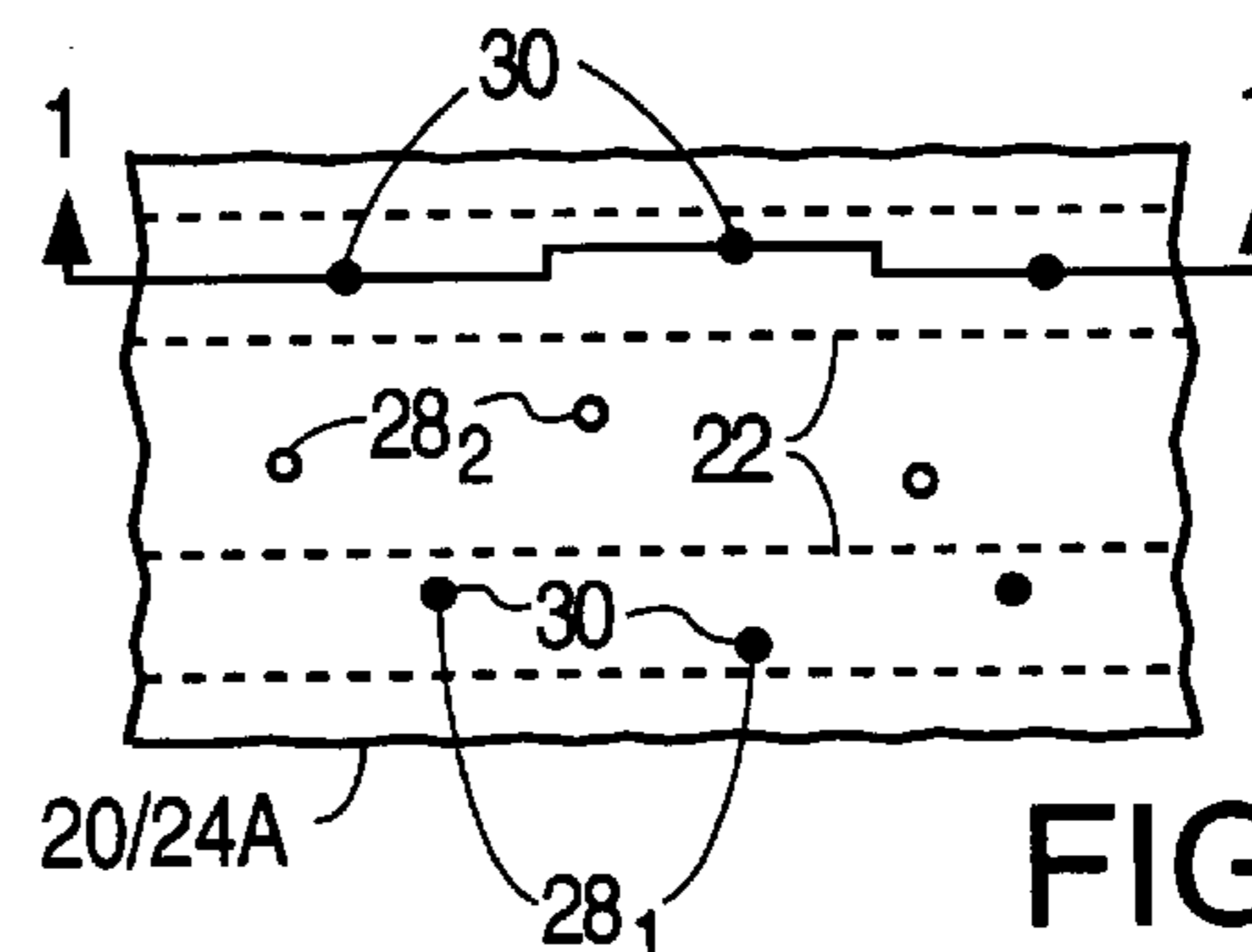


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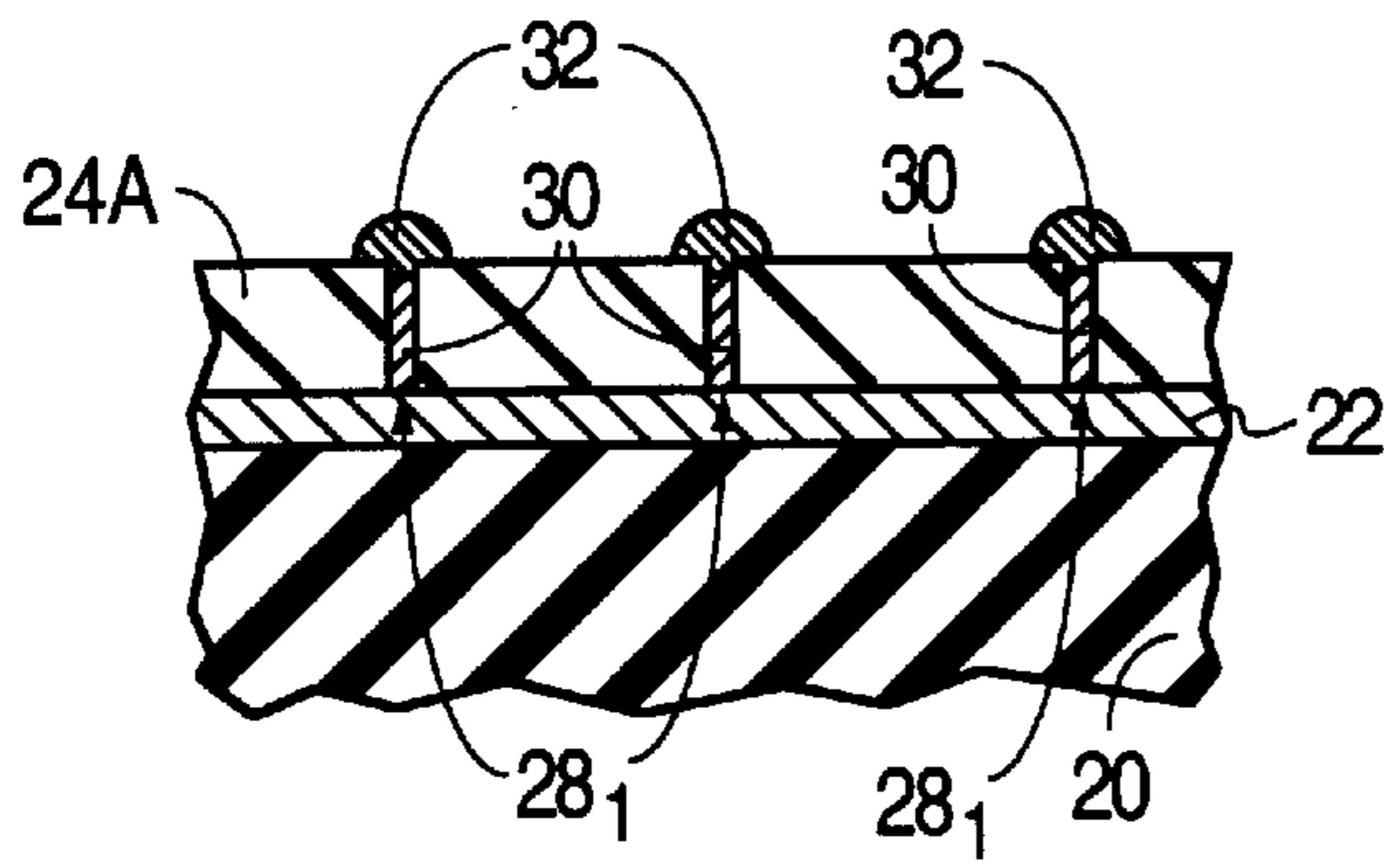


FIG. 1f

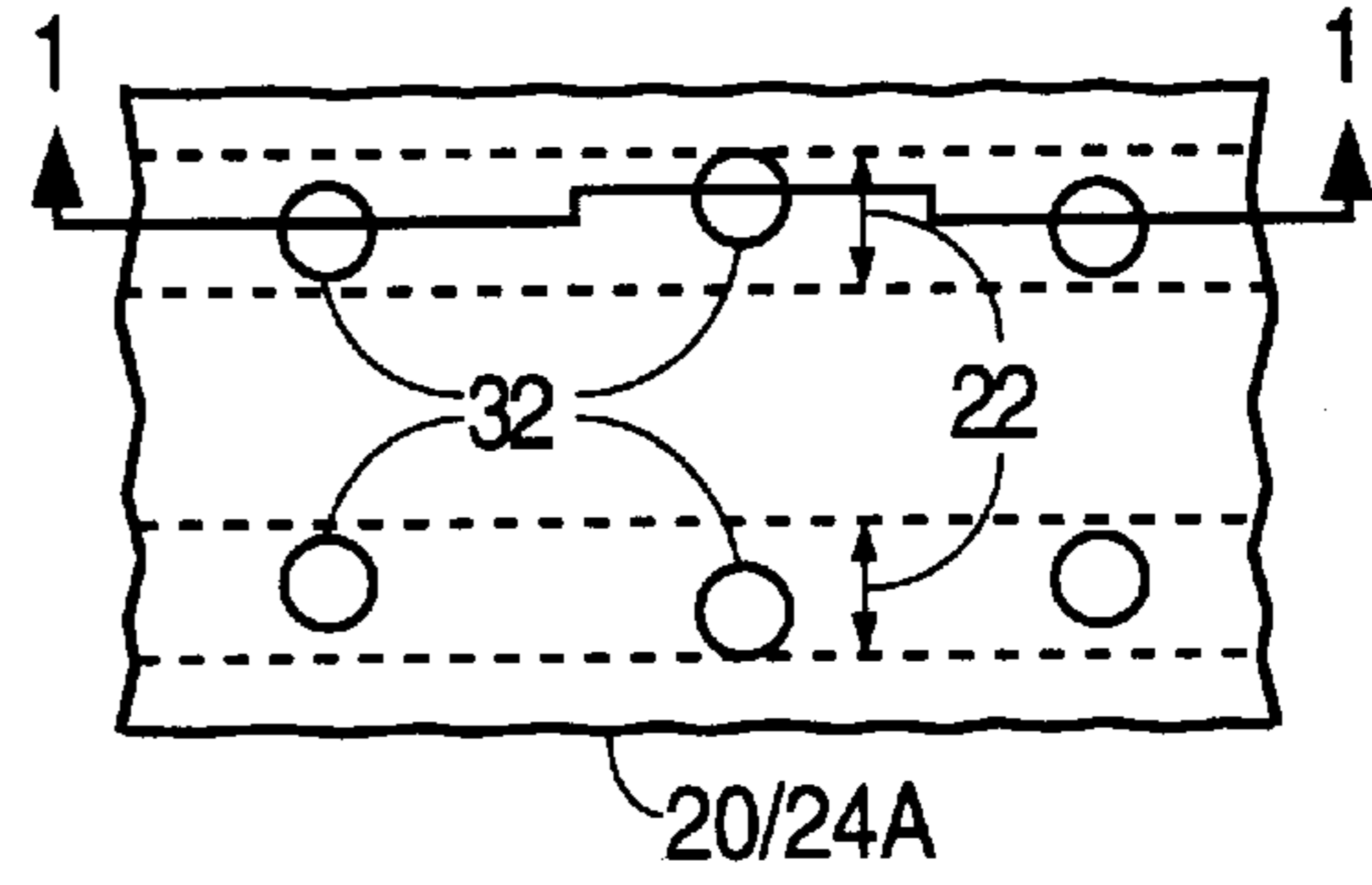


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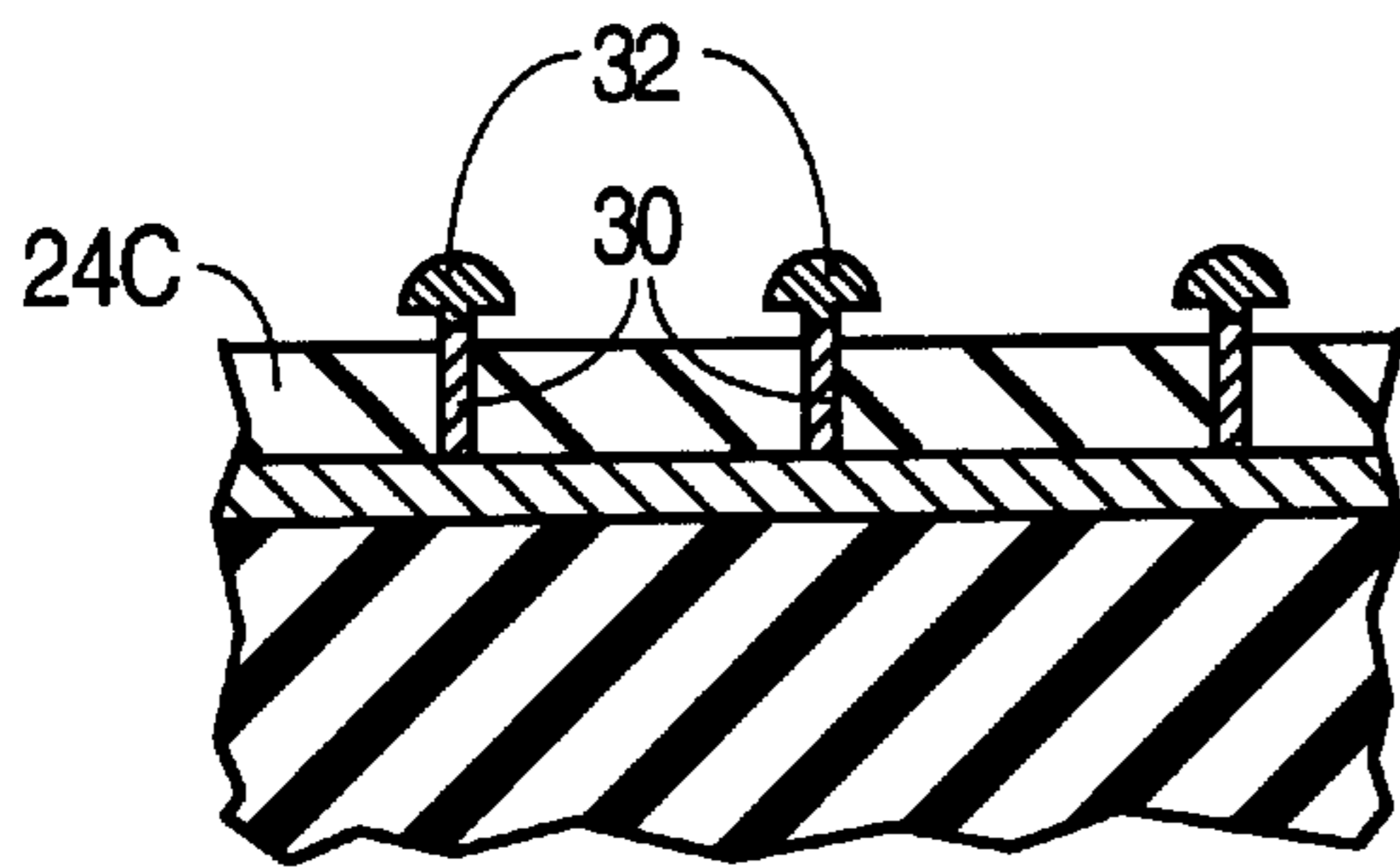


FIG. 1g

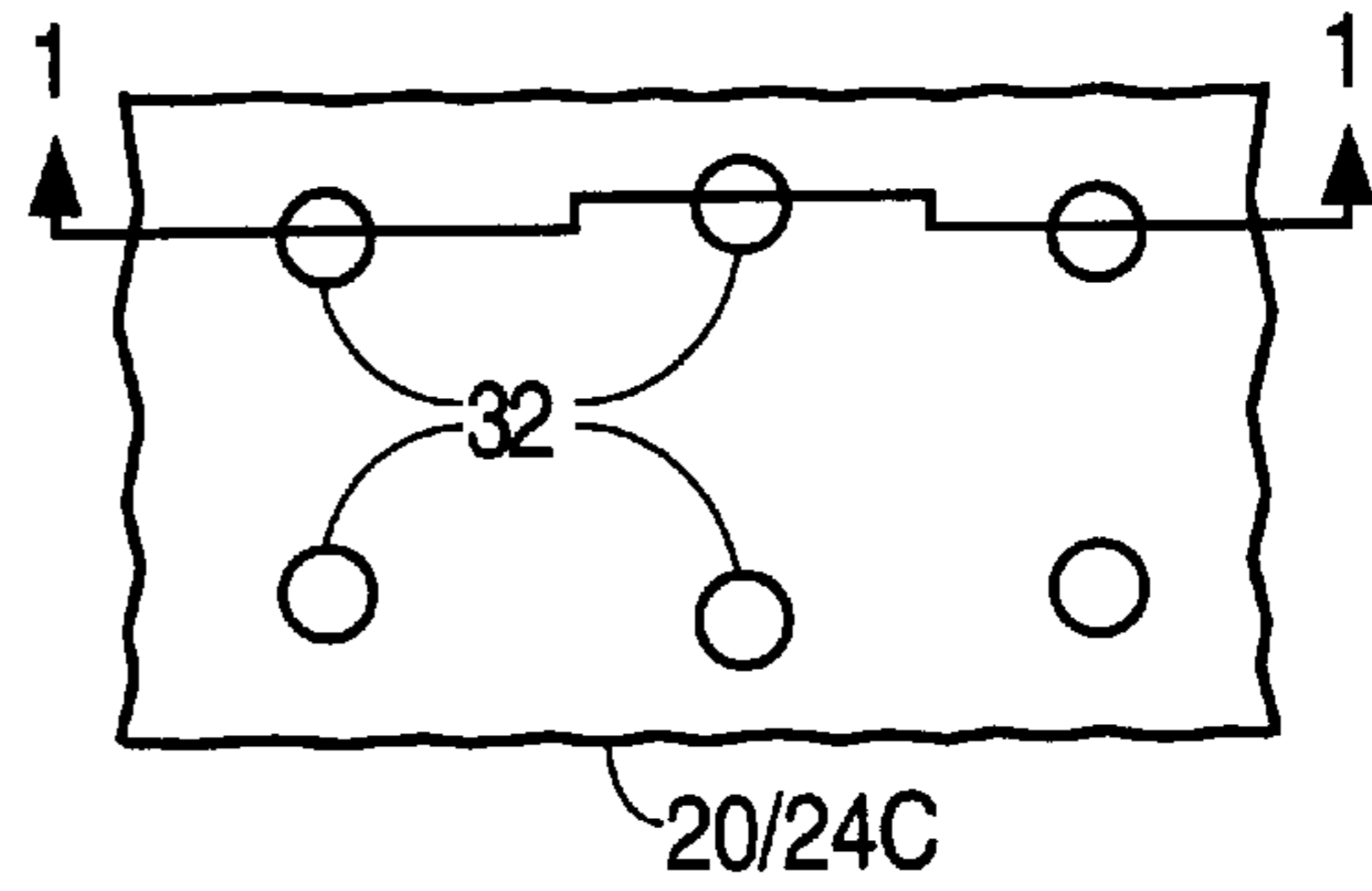


FIG. 2g

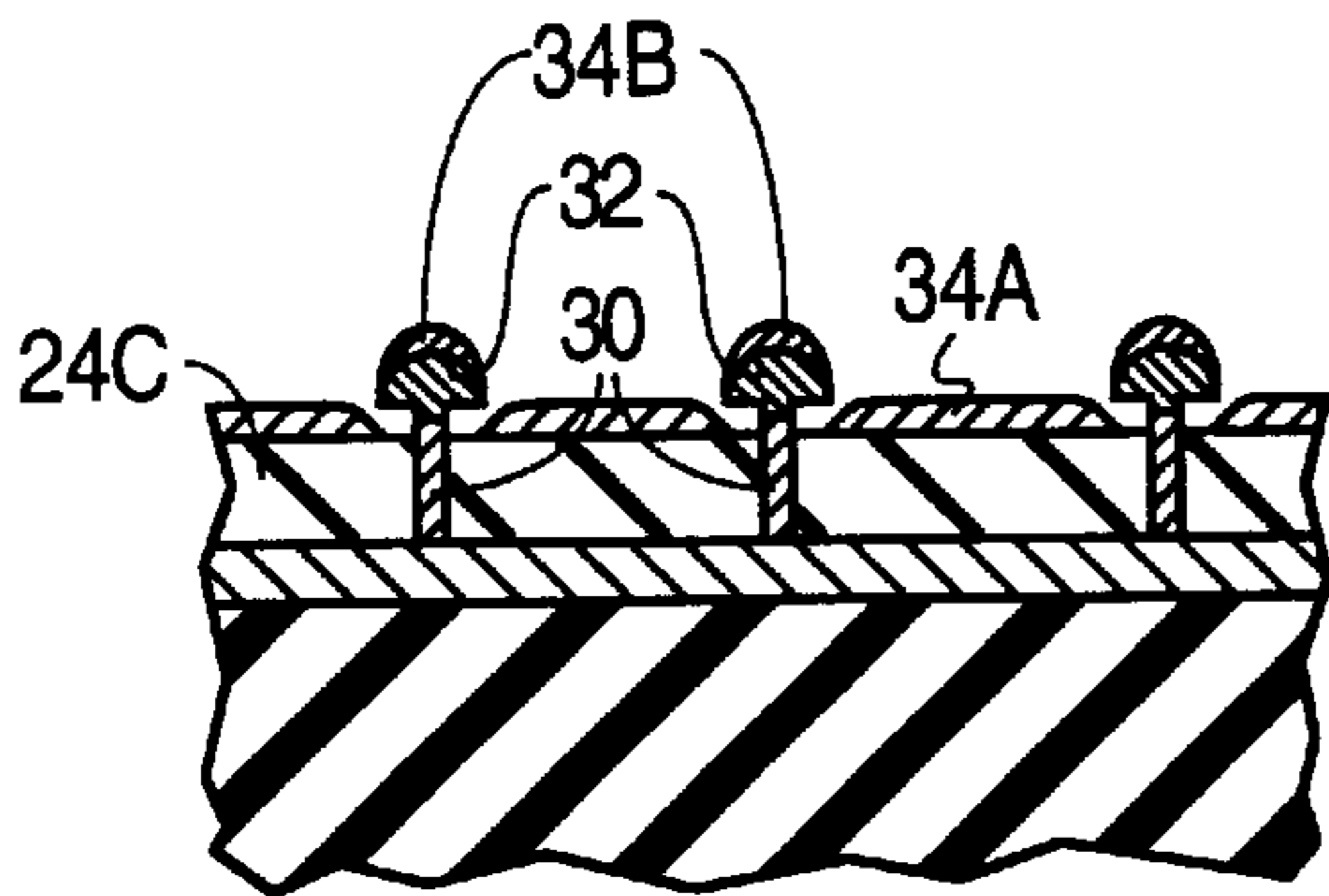


FIG. 1h

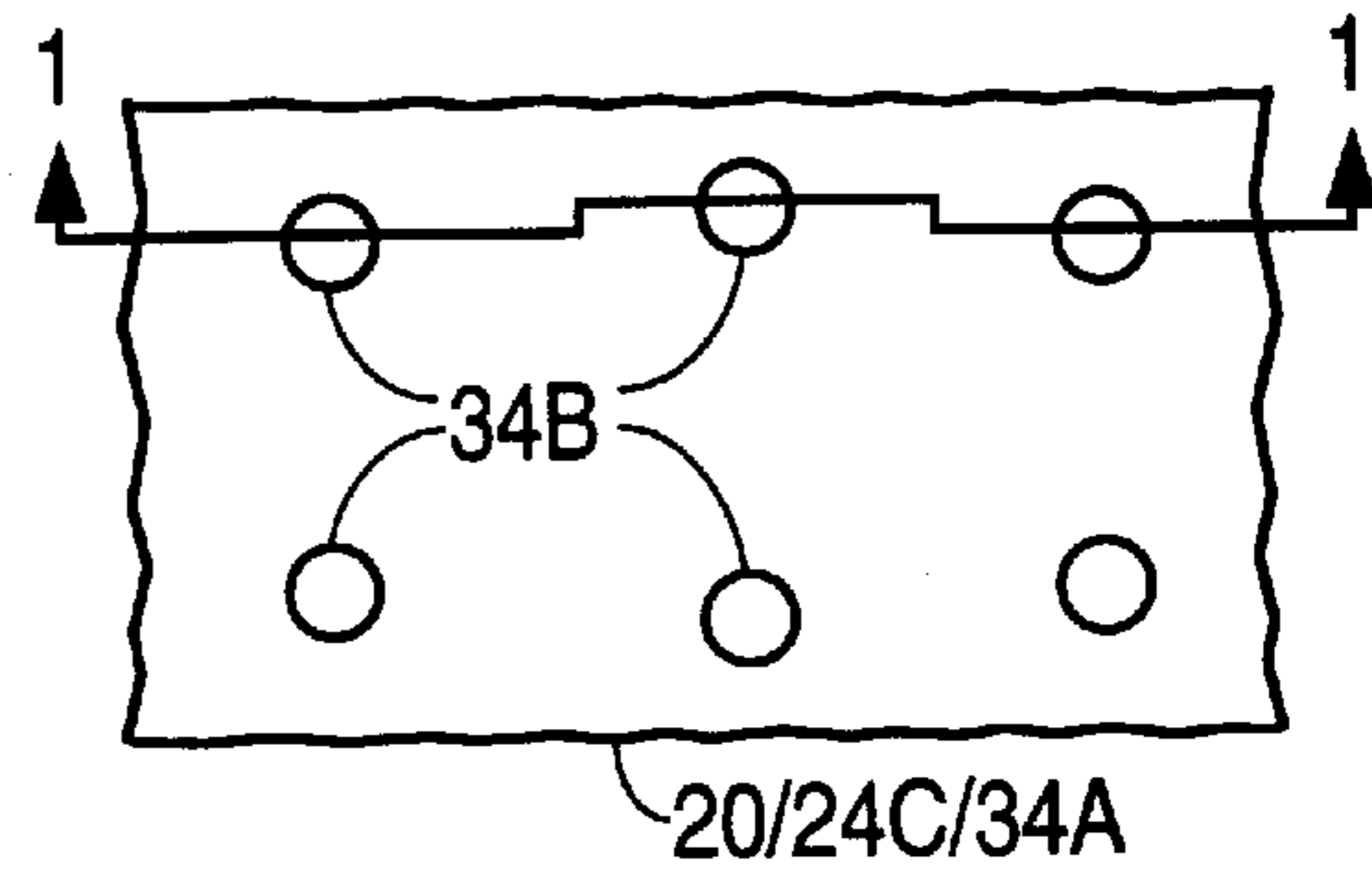


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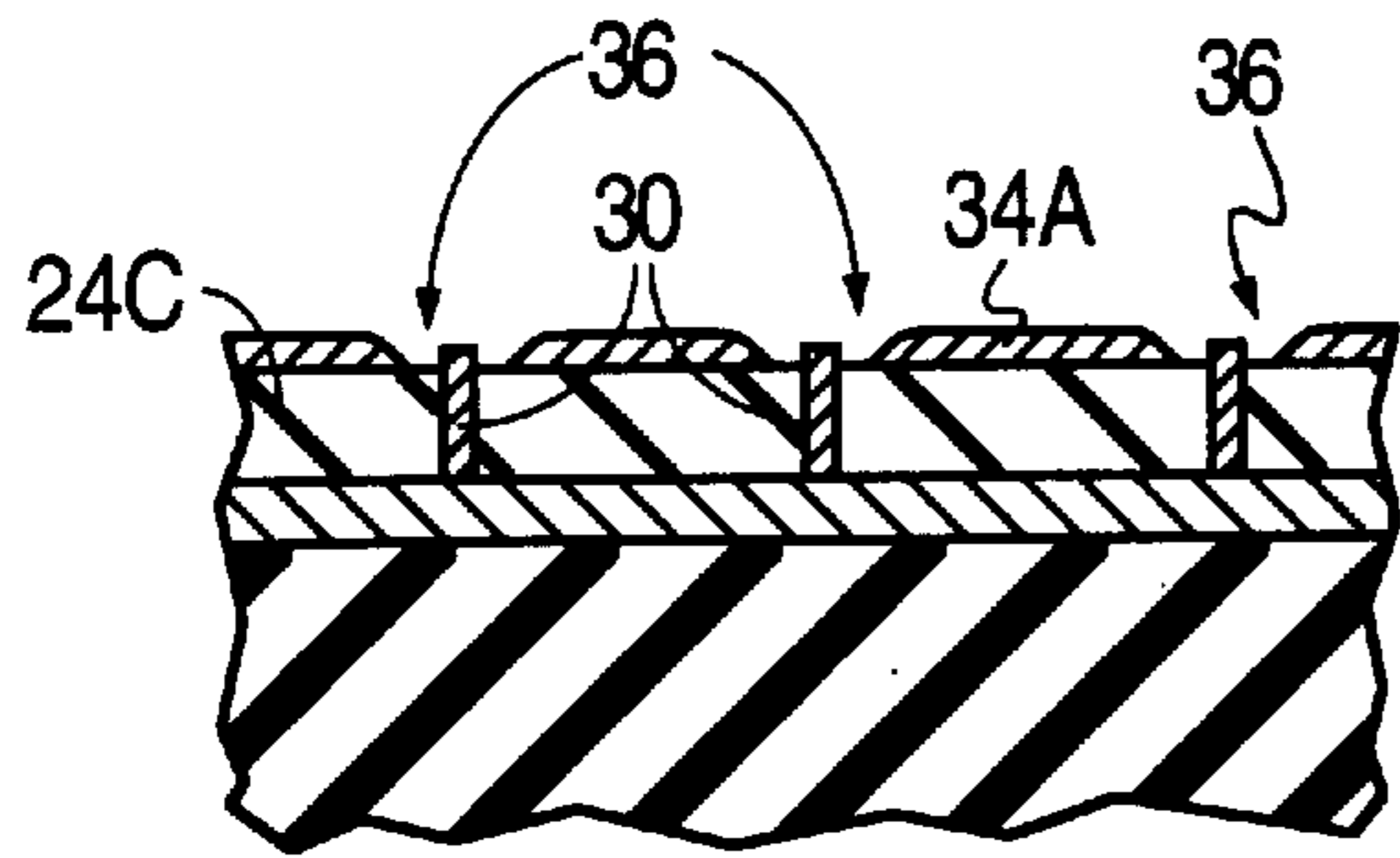


FIG. 1i

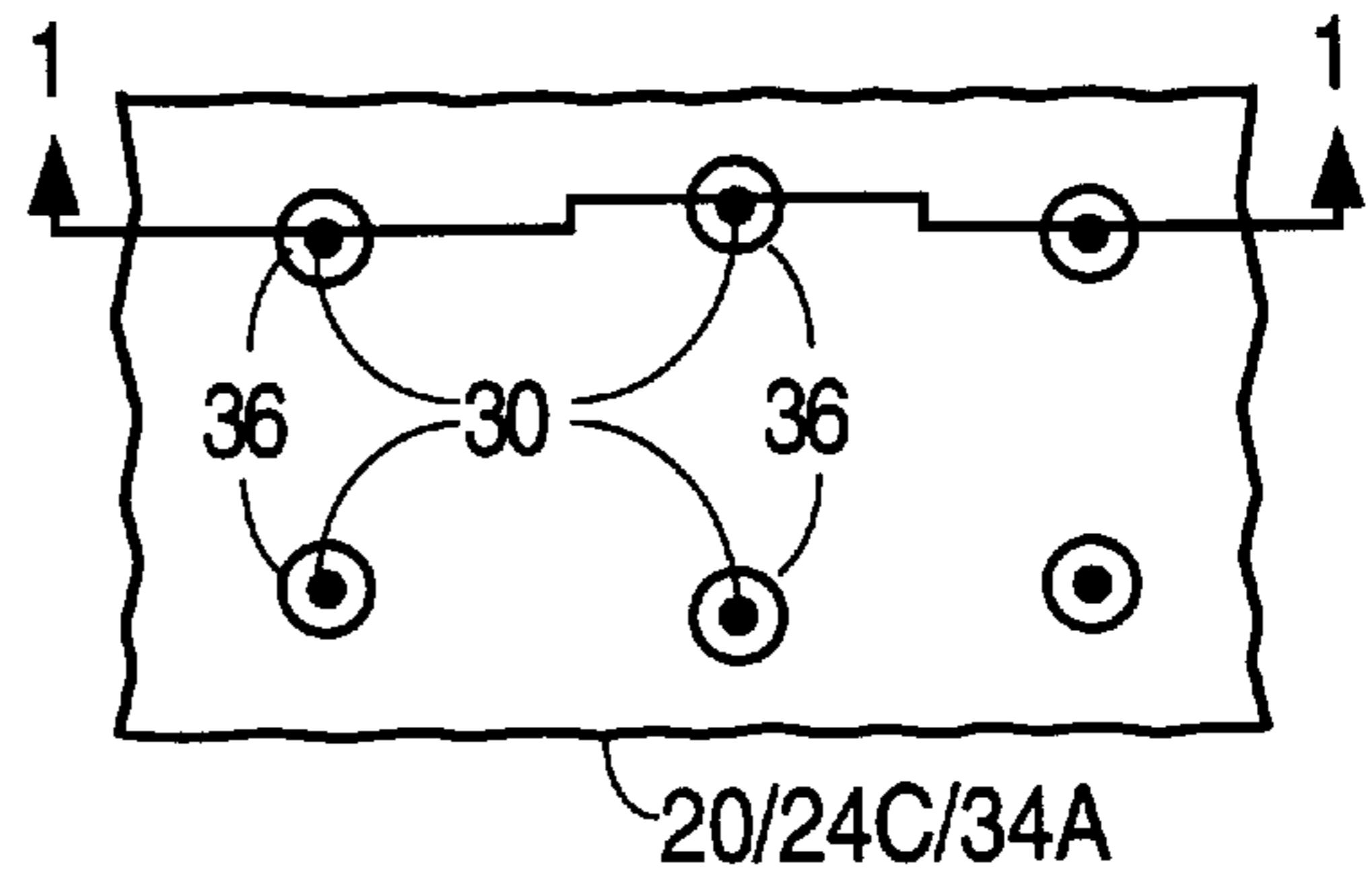


FIG. 2i

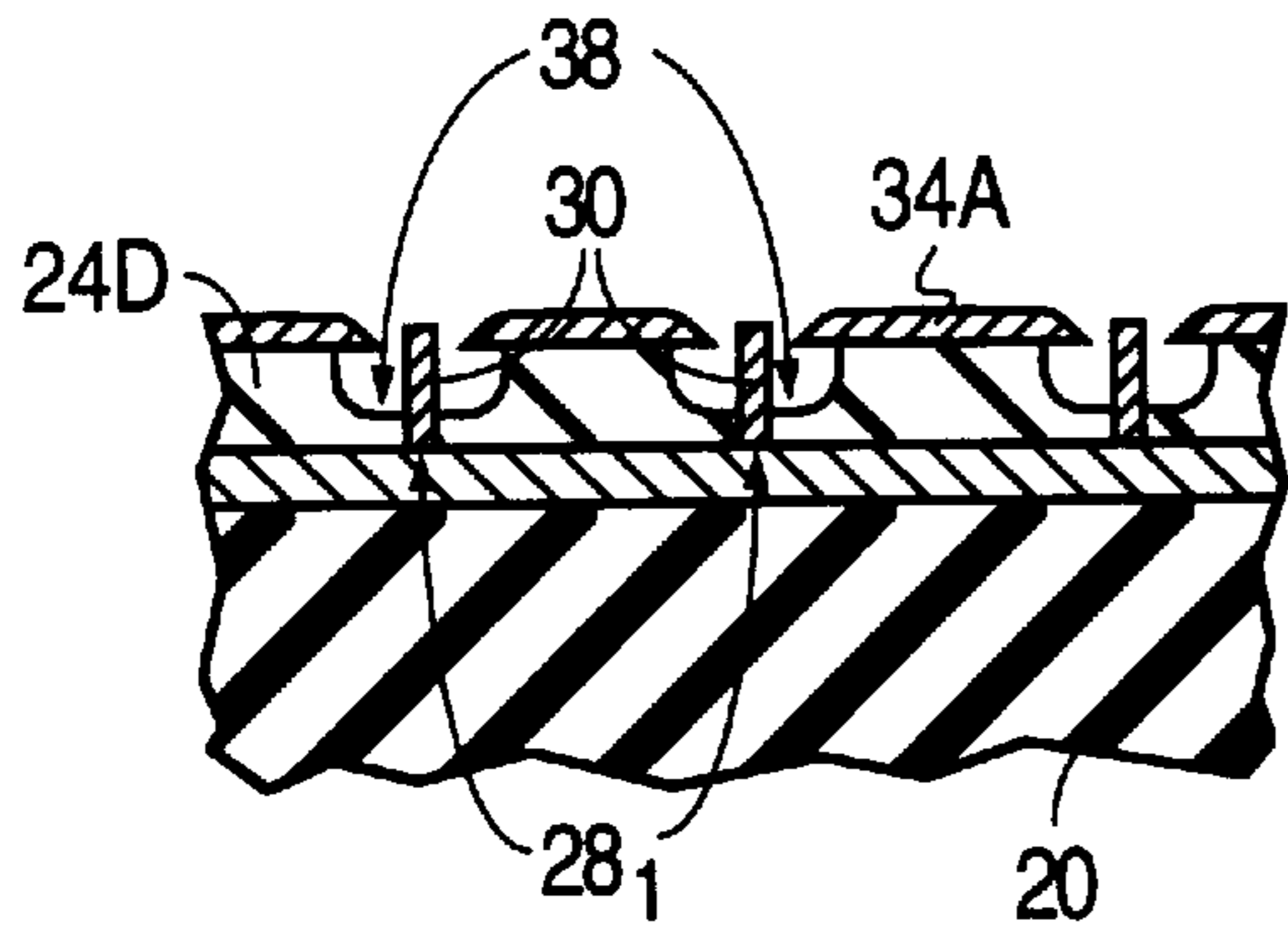


FIG. 1j

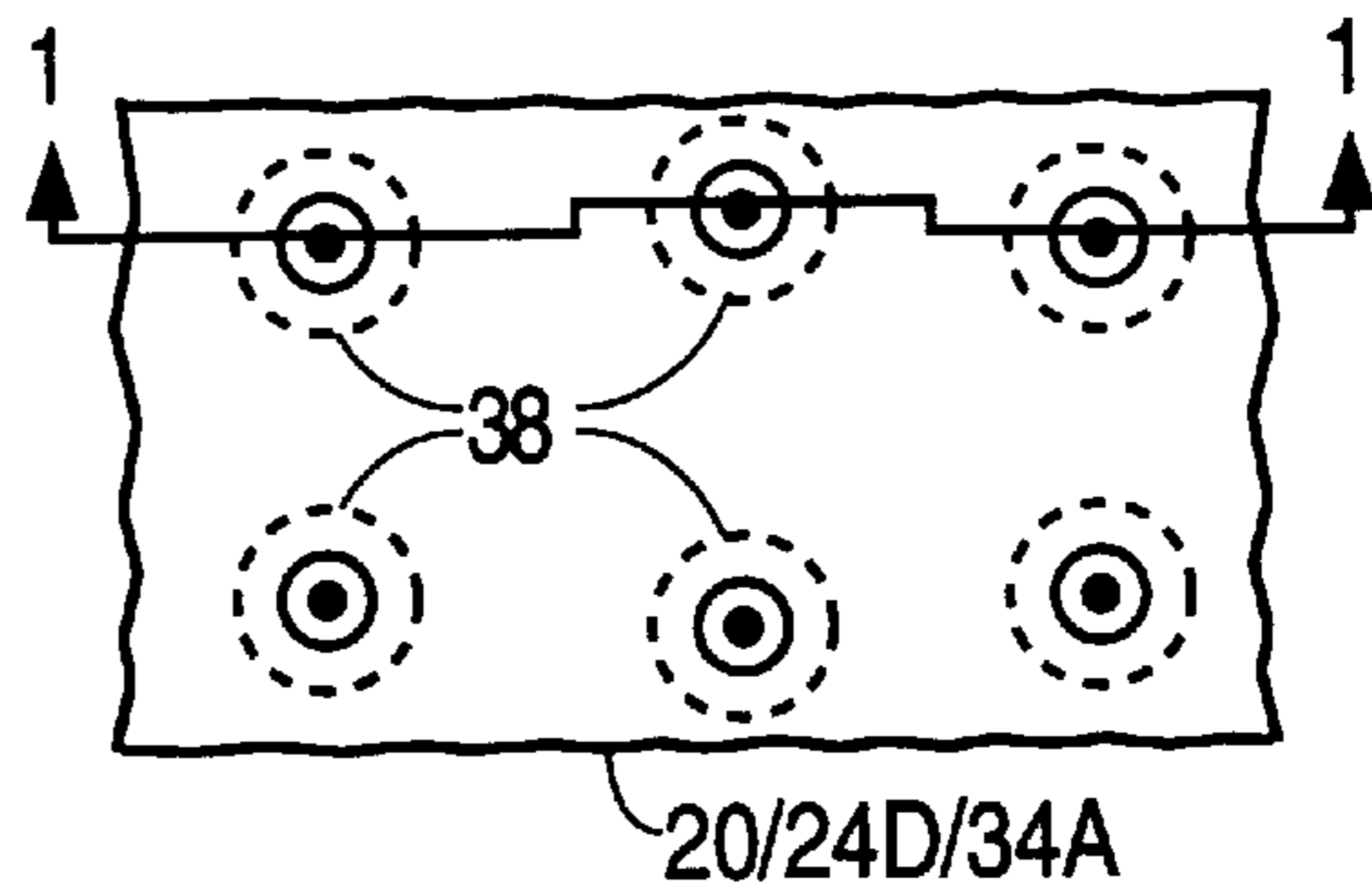


FIG. 2j

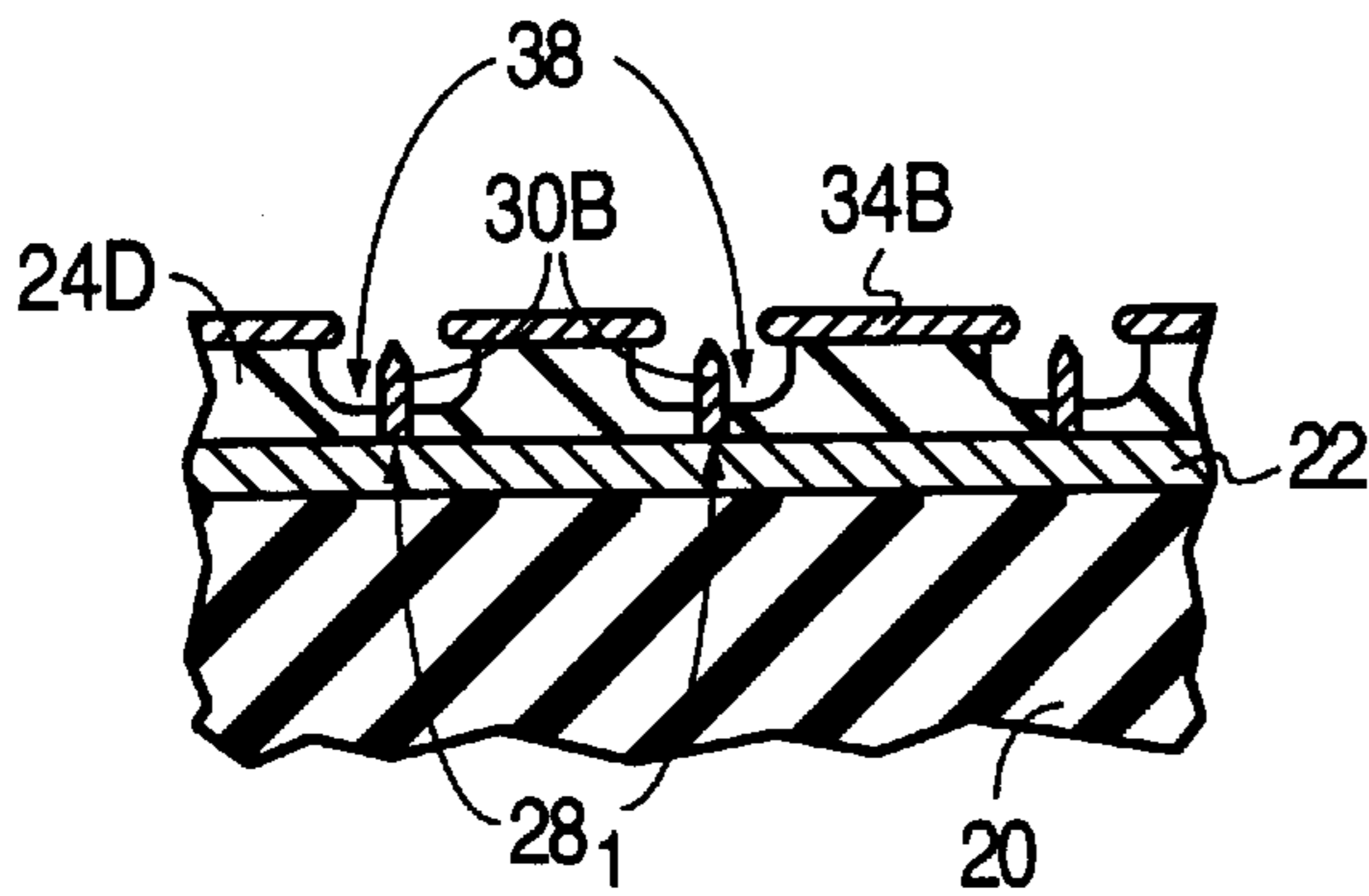


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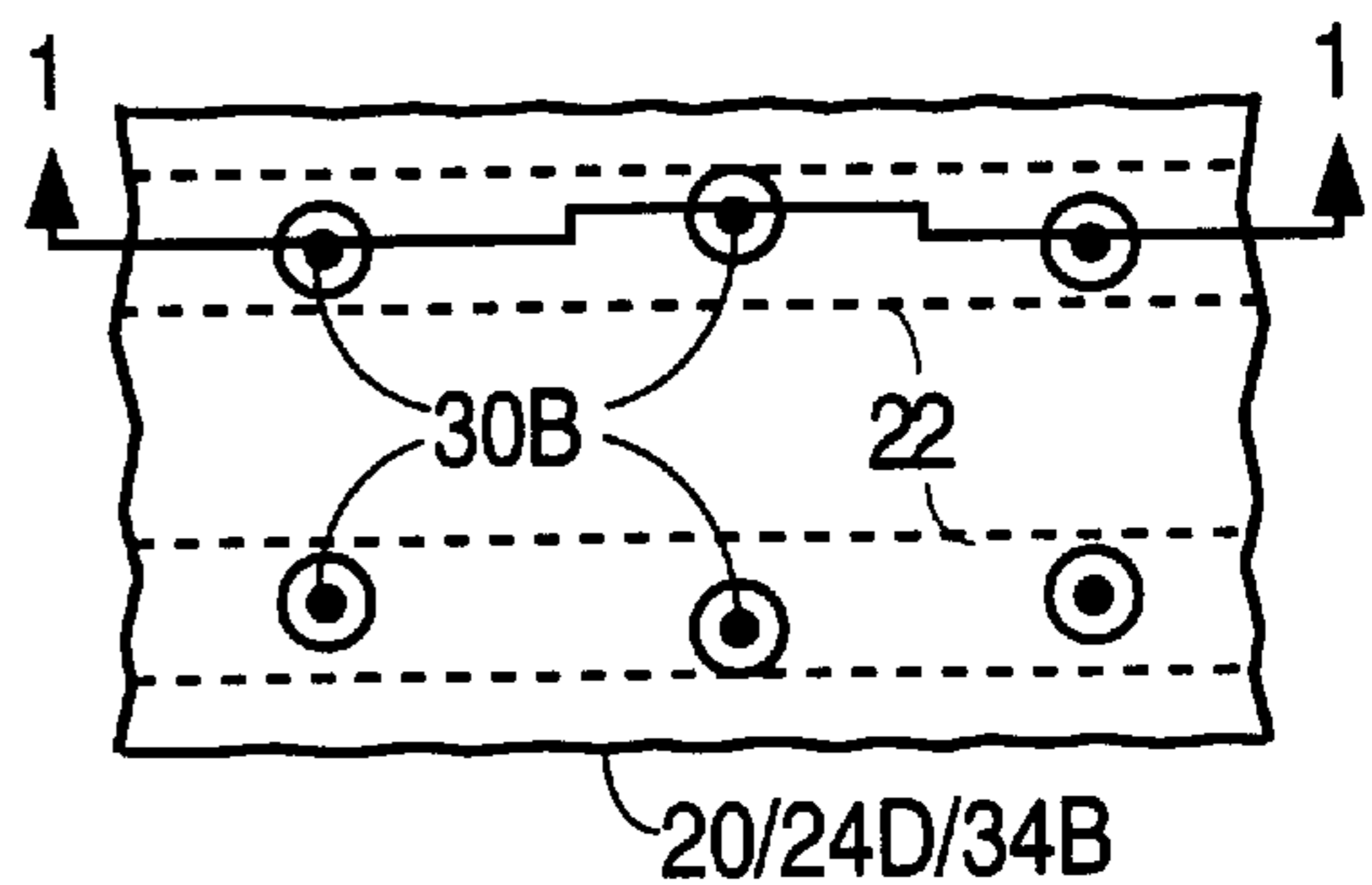


FIG. 2k

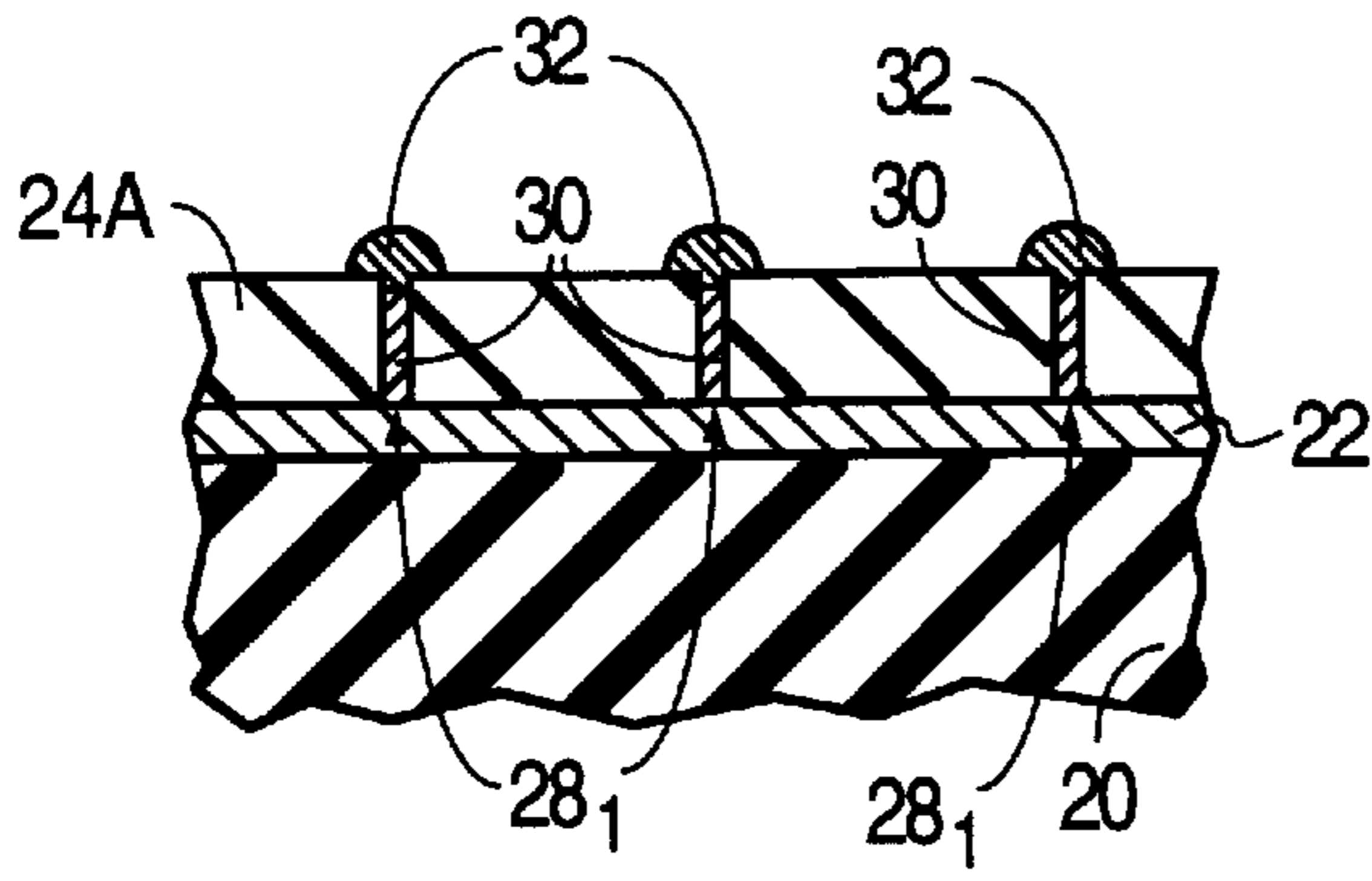


FIG. 3a

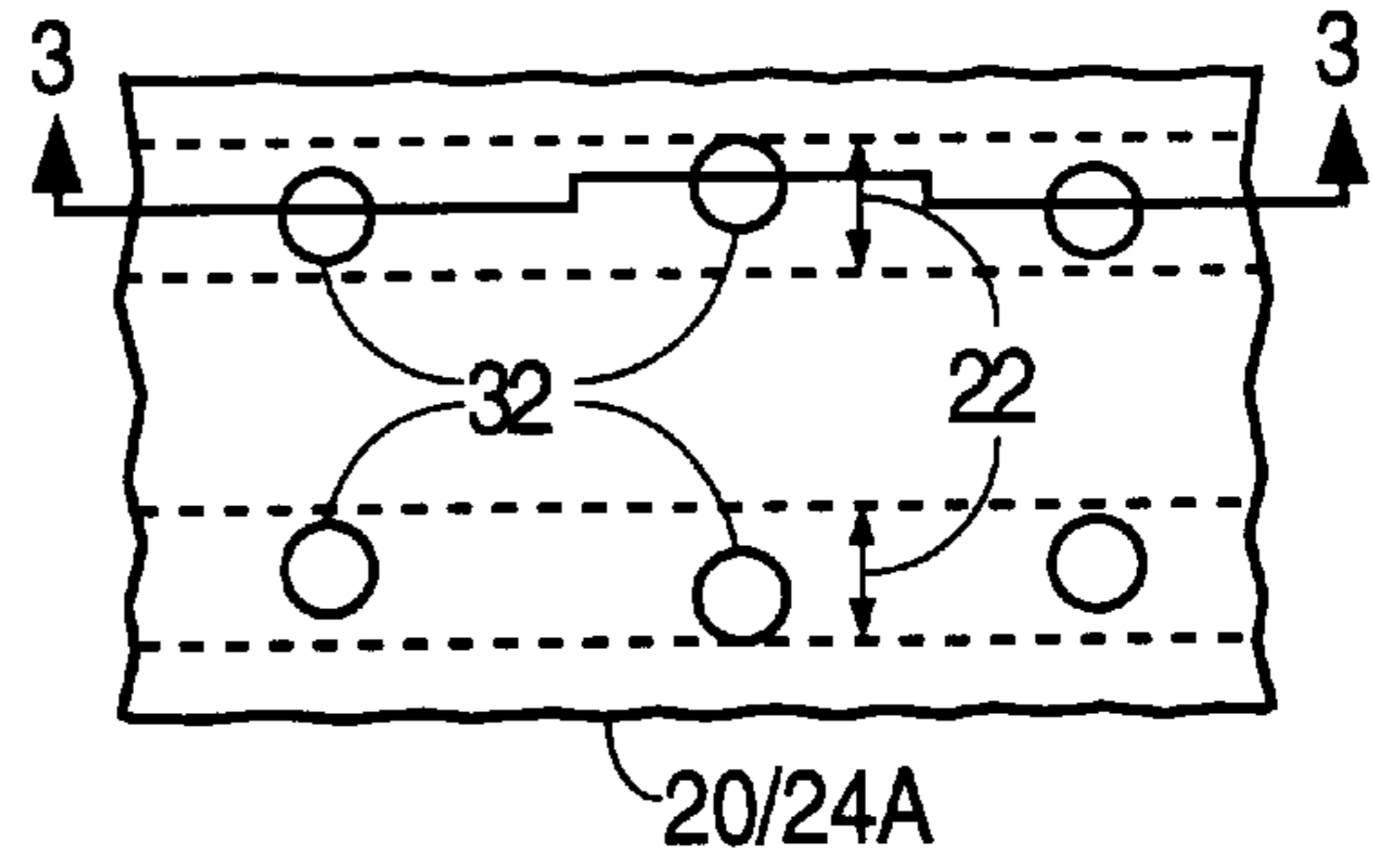


FIG. 4a

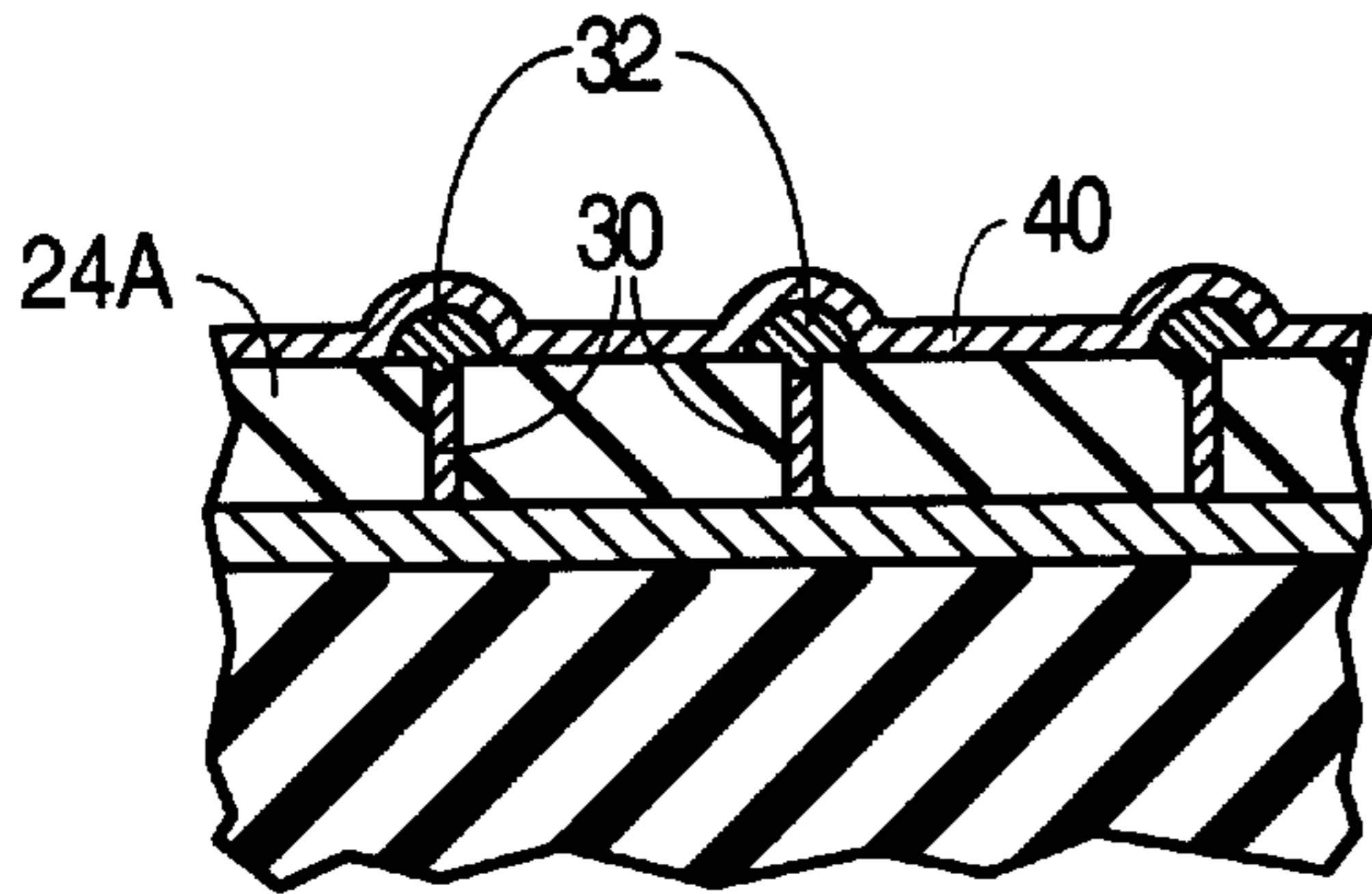


FIG. 3b

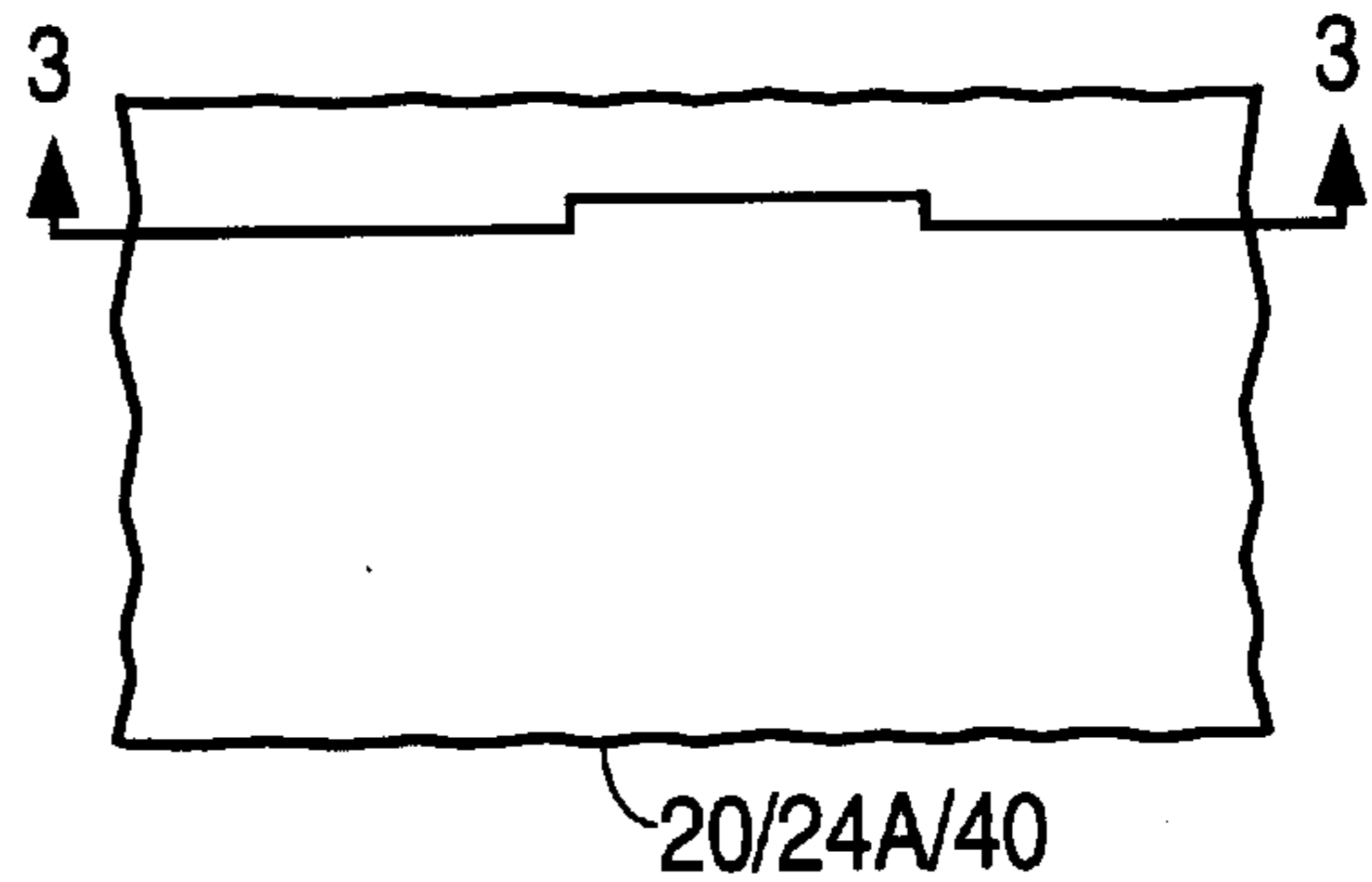


FIG. 4b

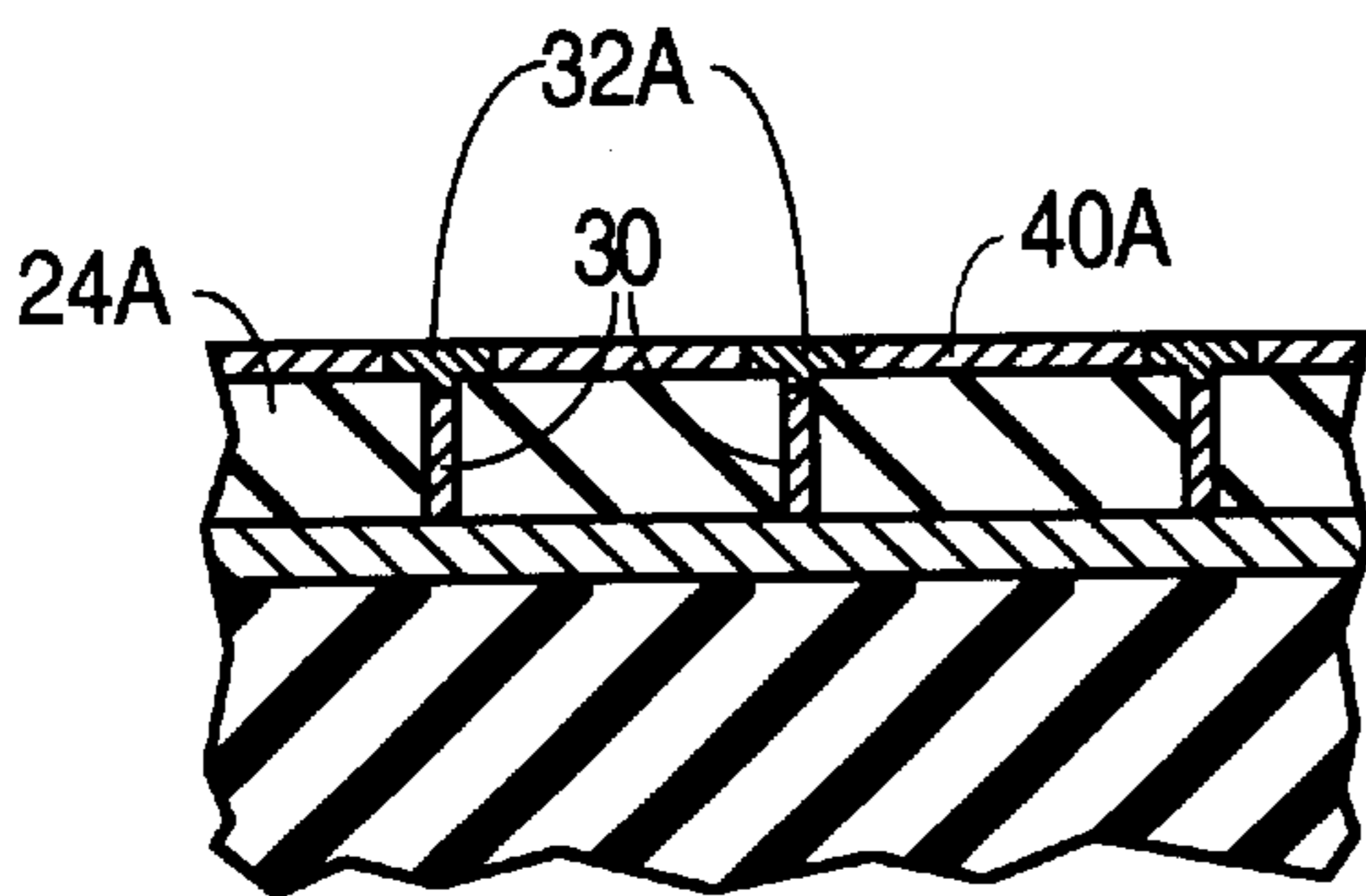


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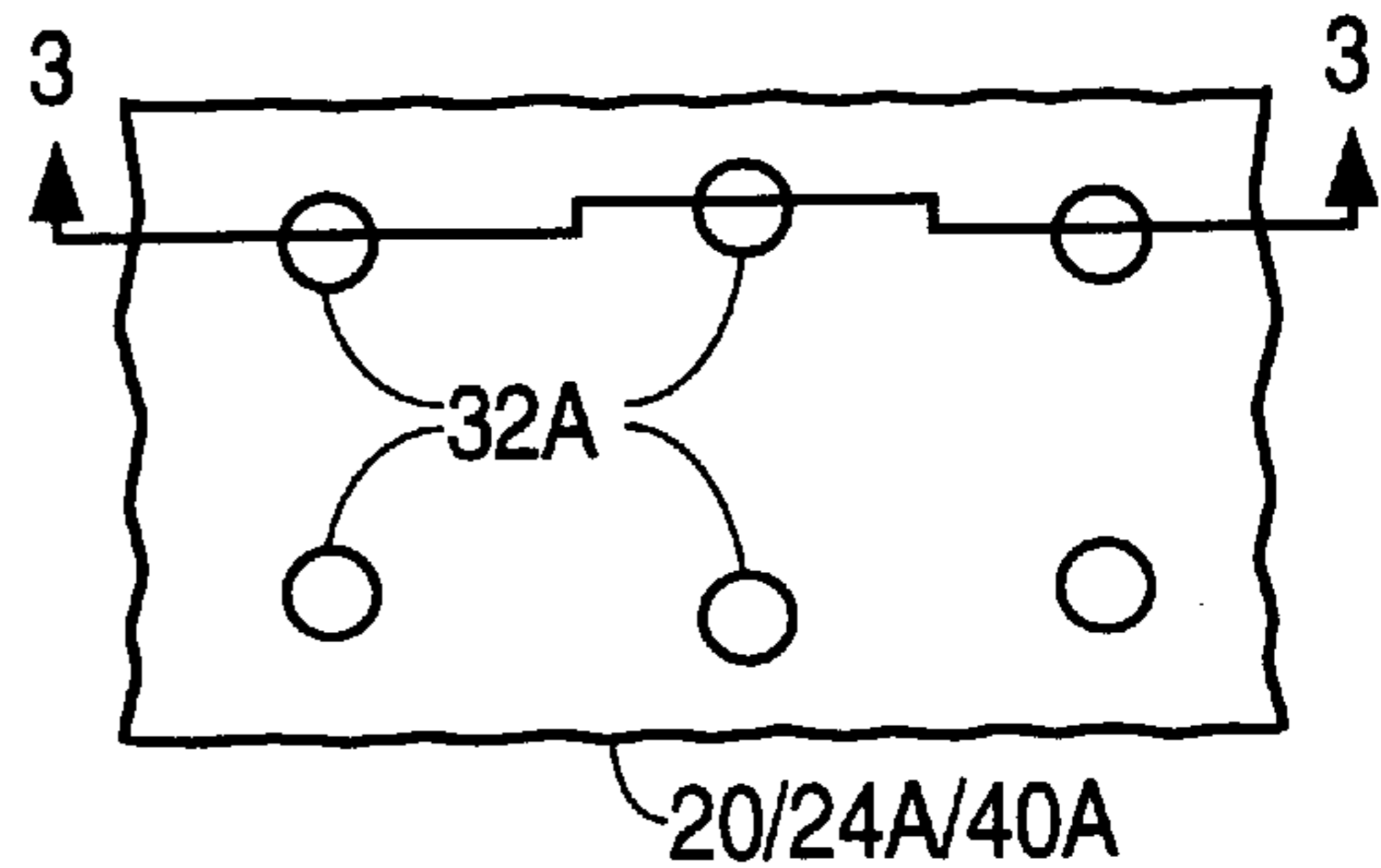


FIG. 4c

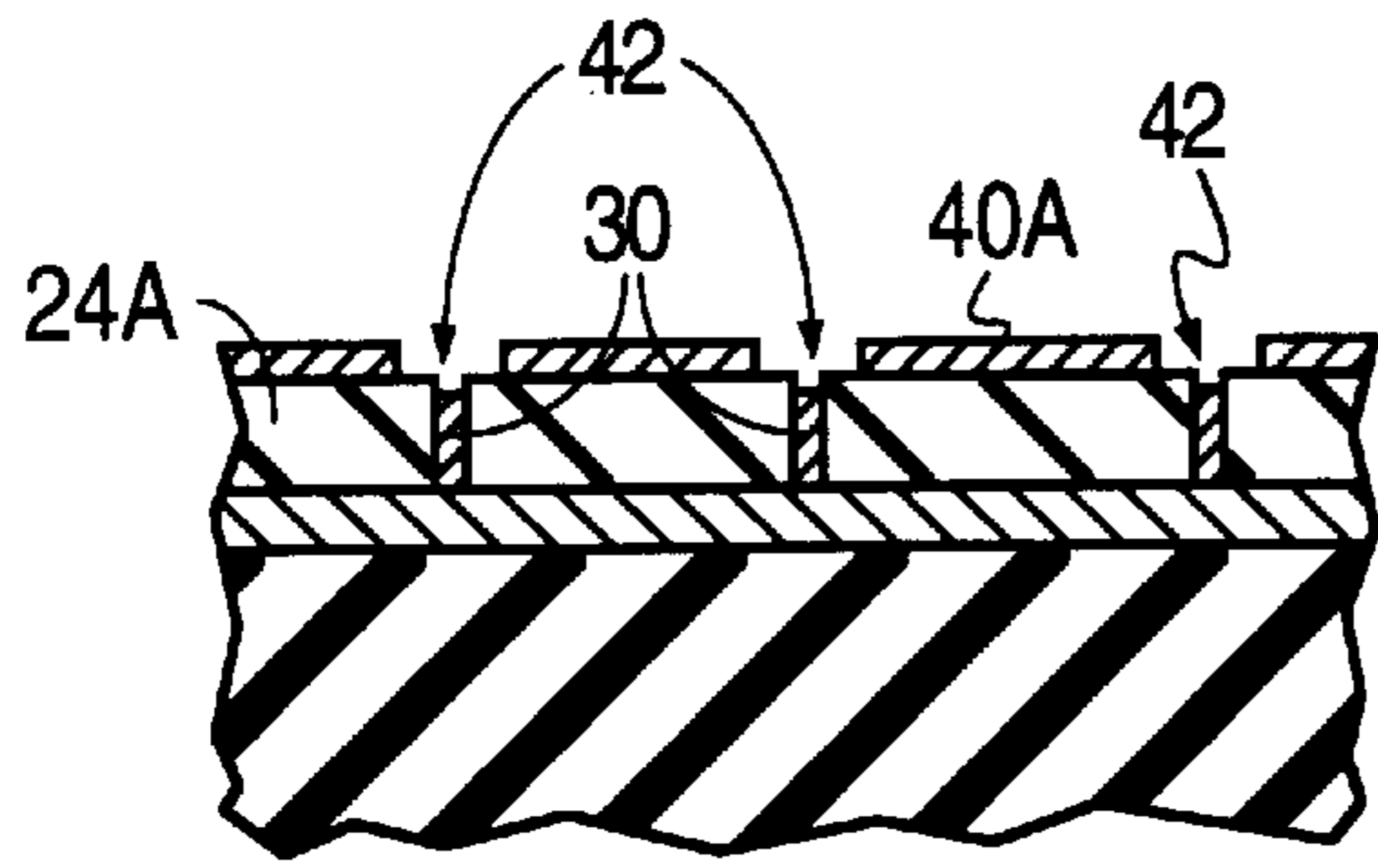


FIG. 3d

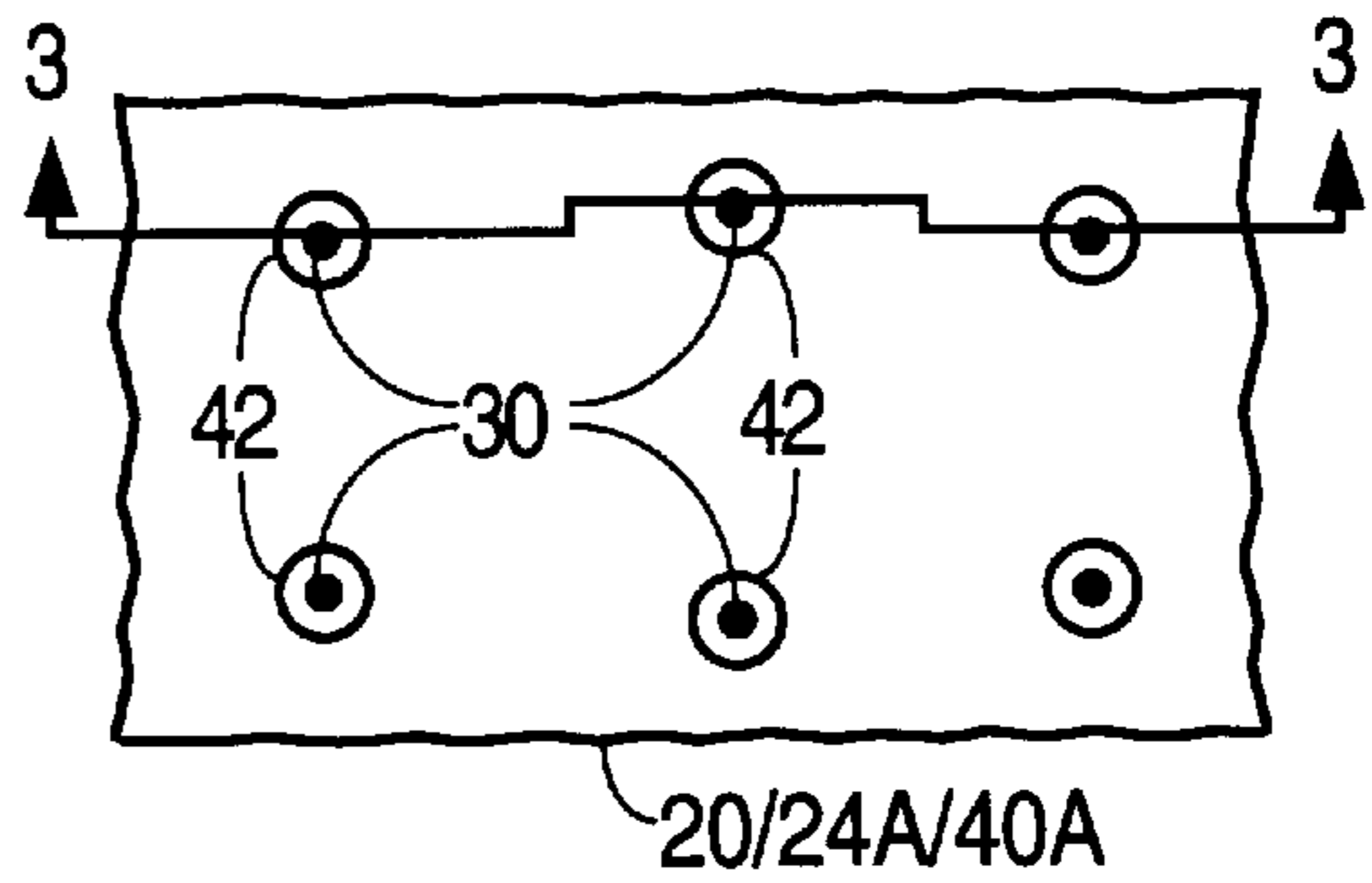


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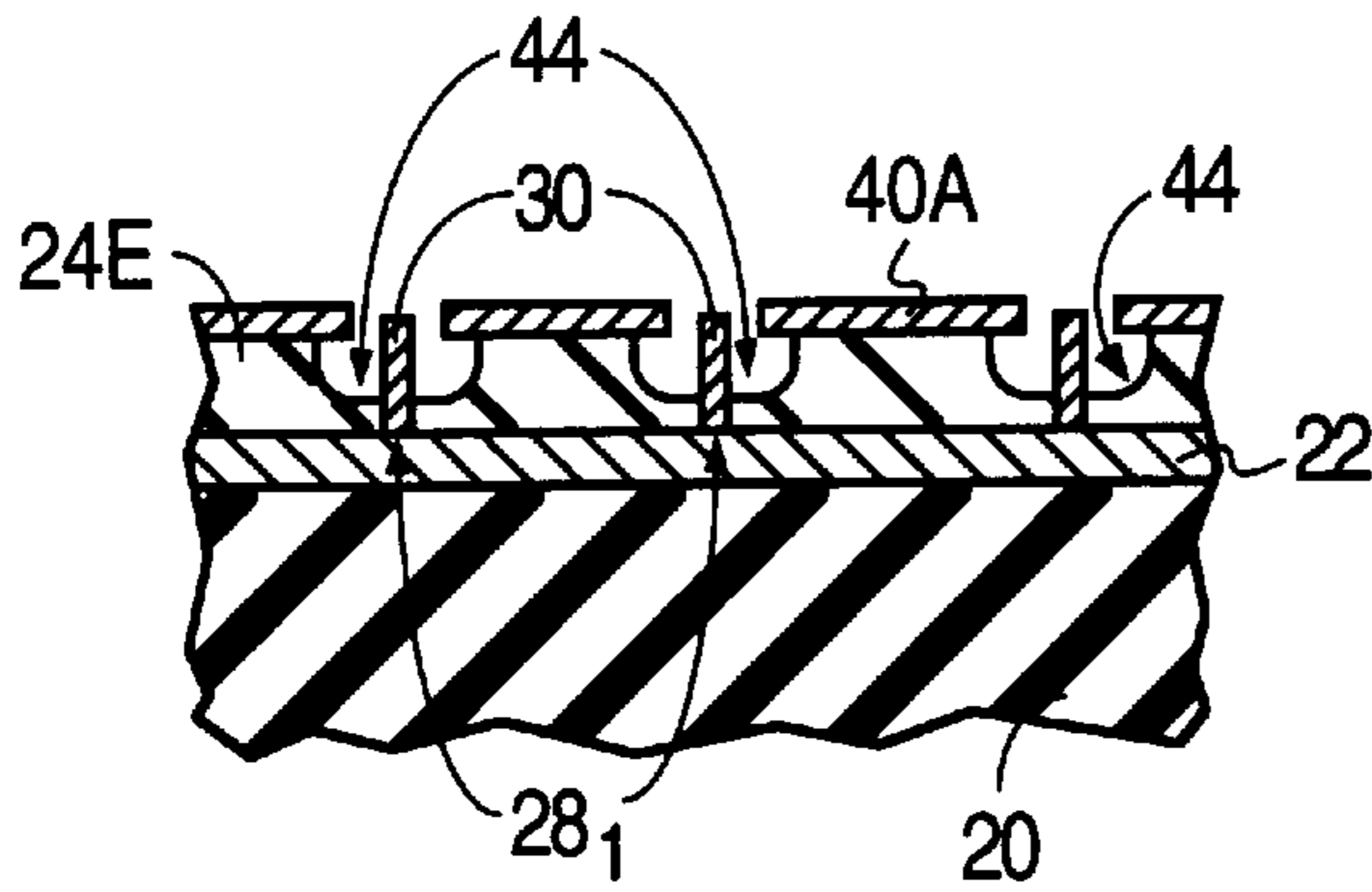


FIG. 3e

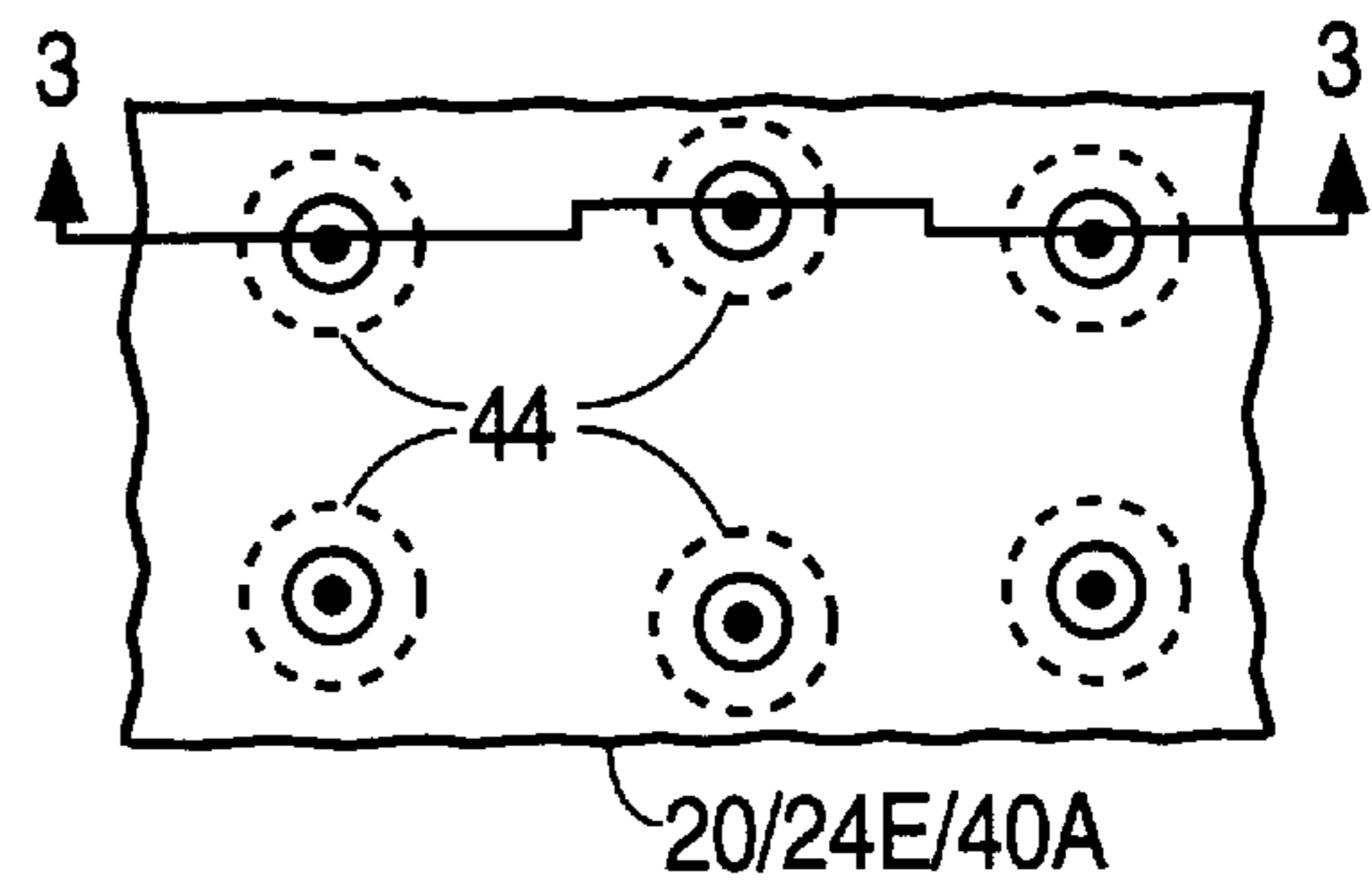


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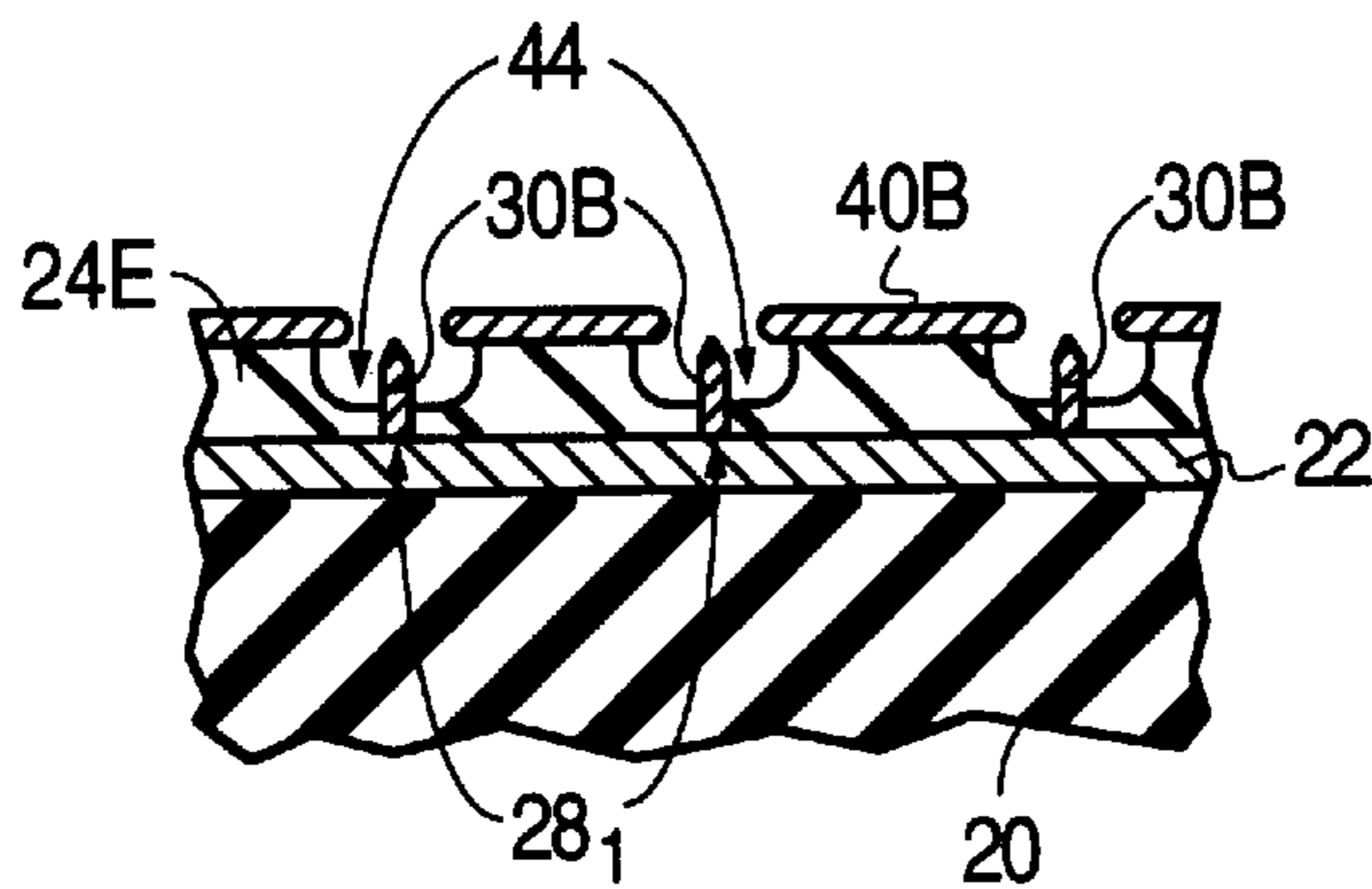


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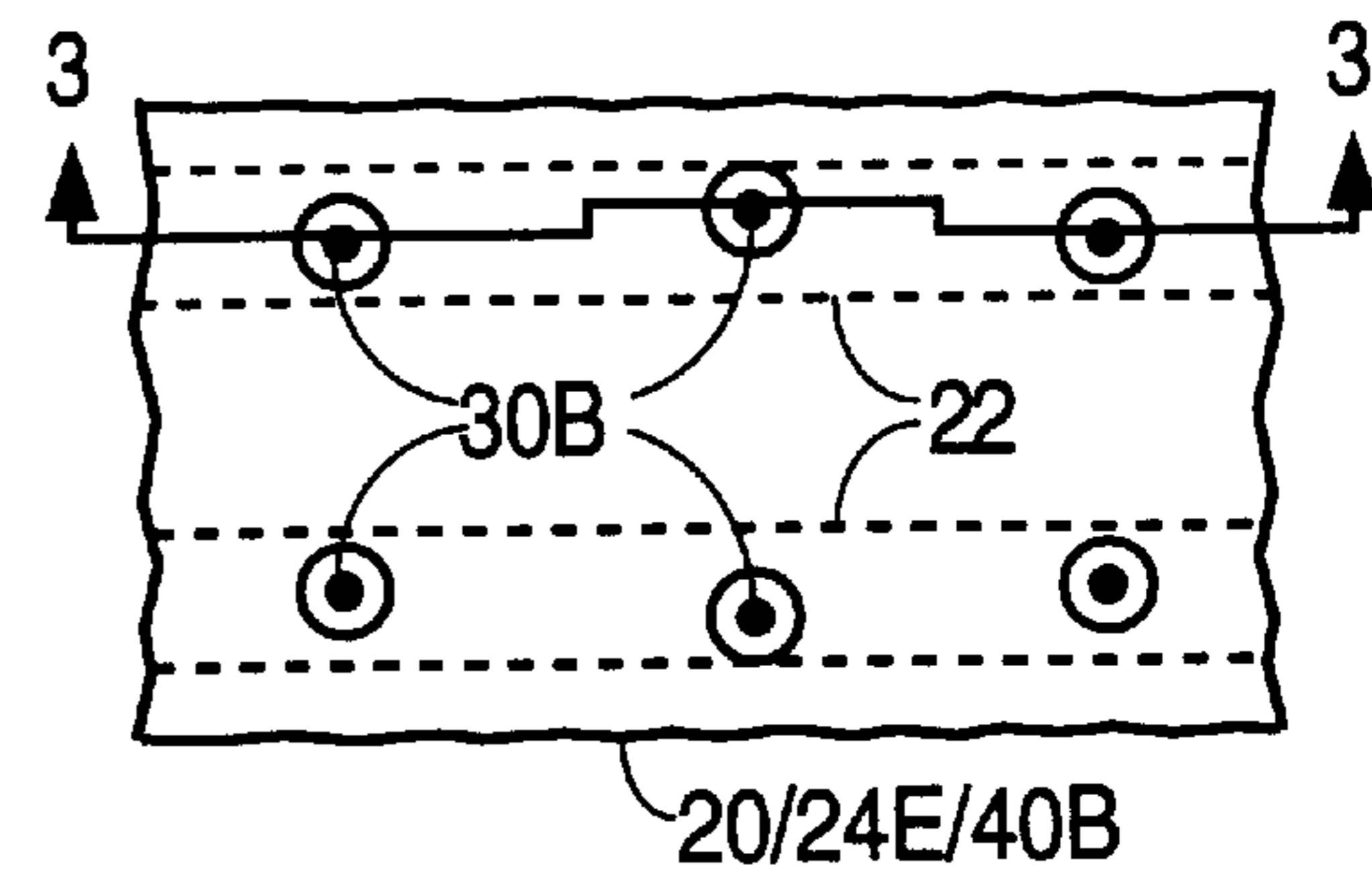


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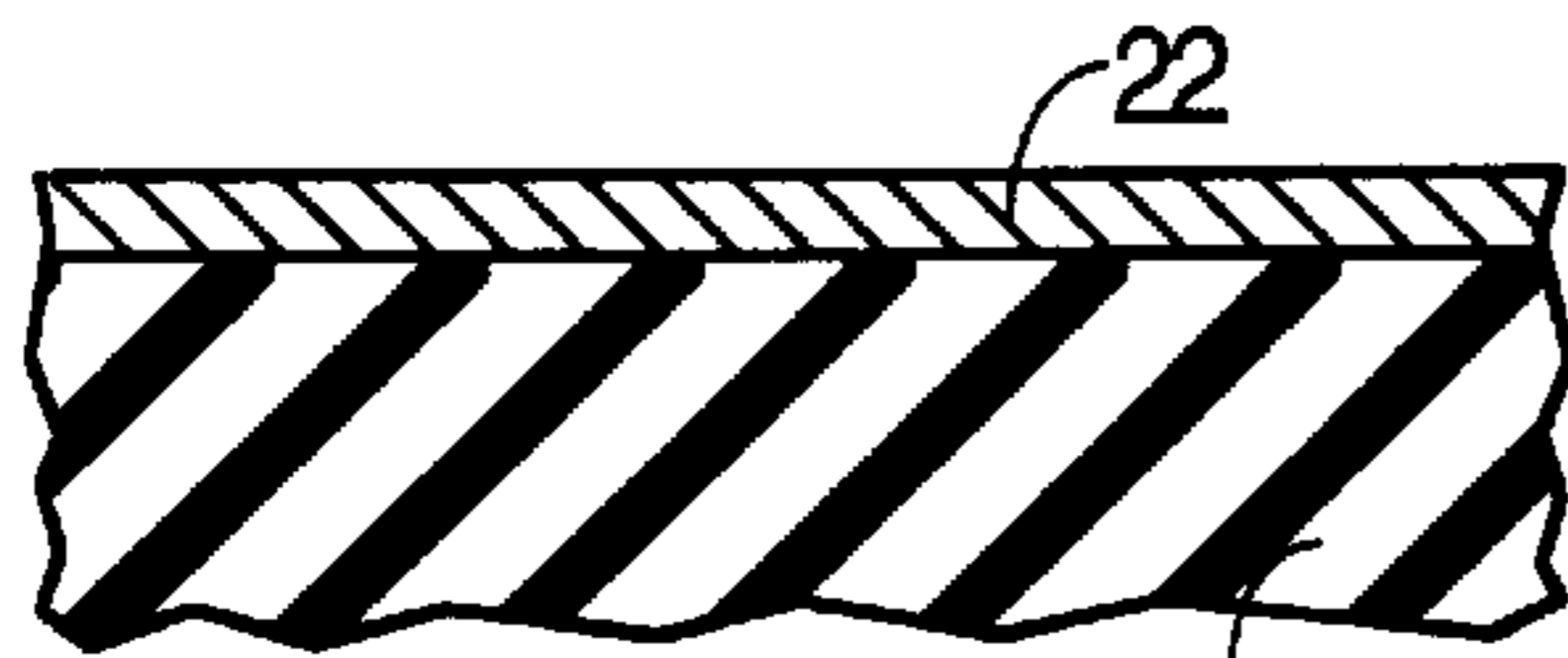


FIG. 5a

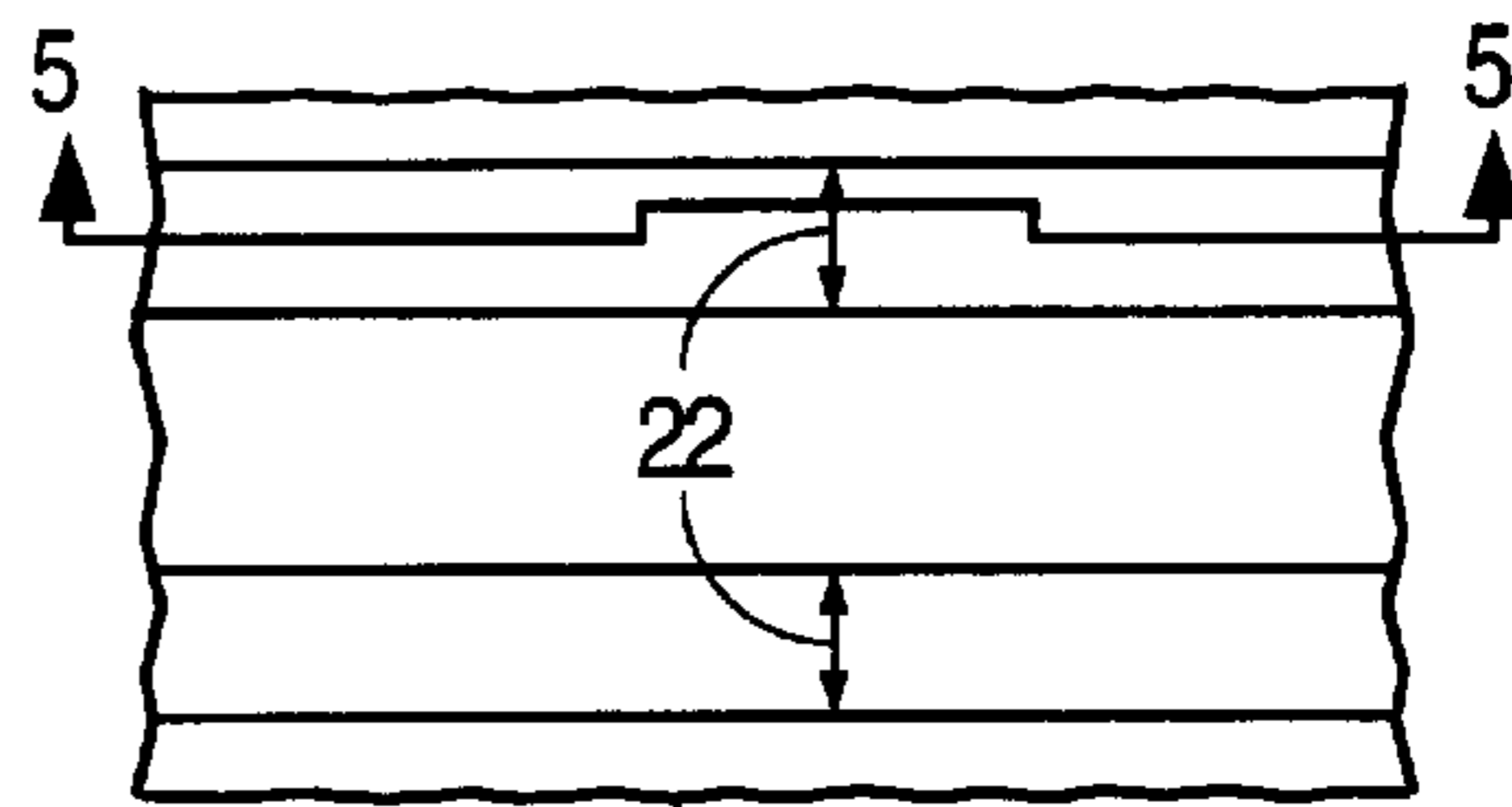


FIG. 6a

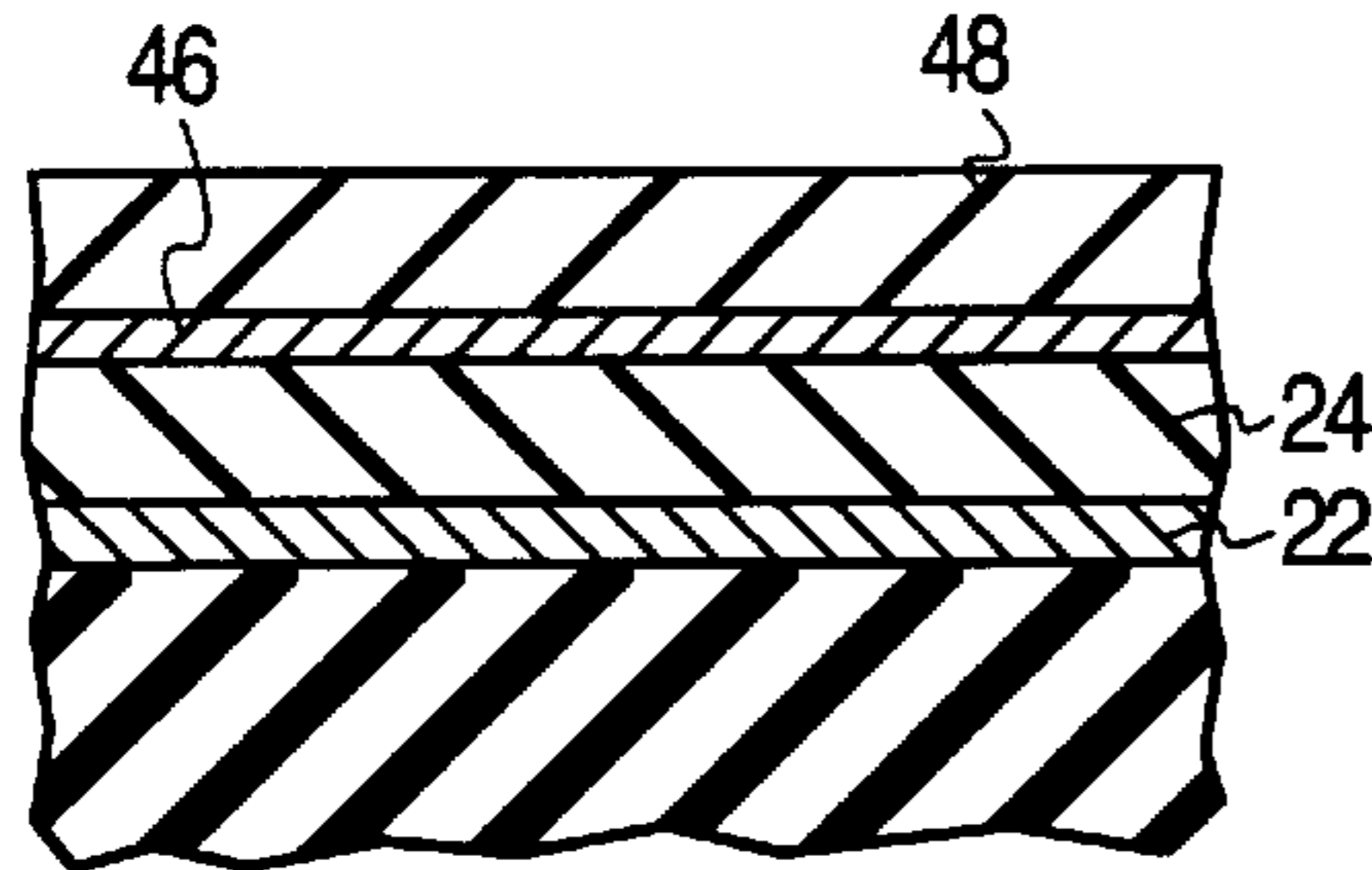


FIG. 5b

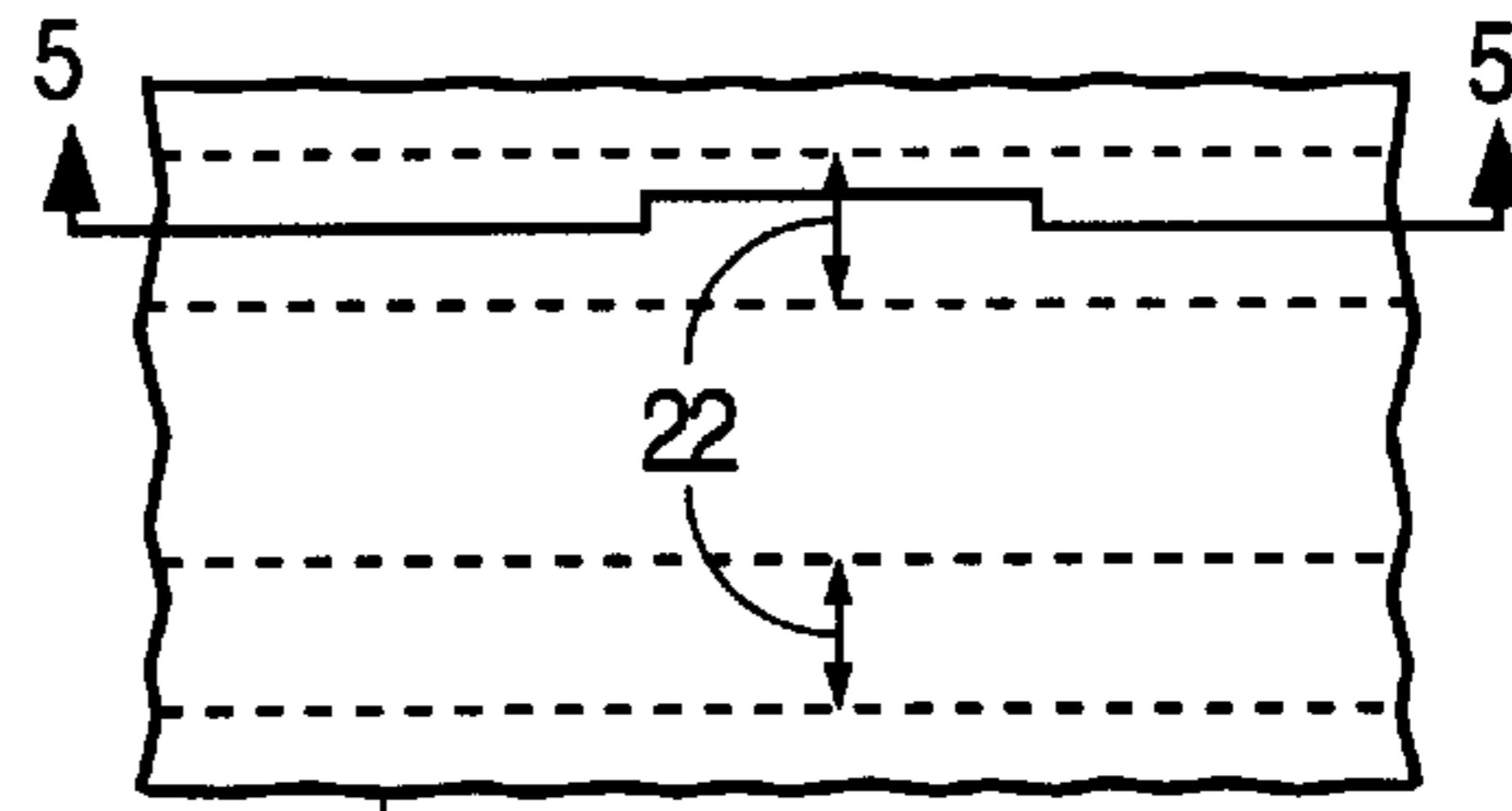


FIG. 6b

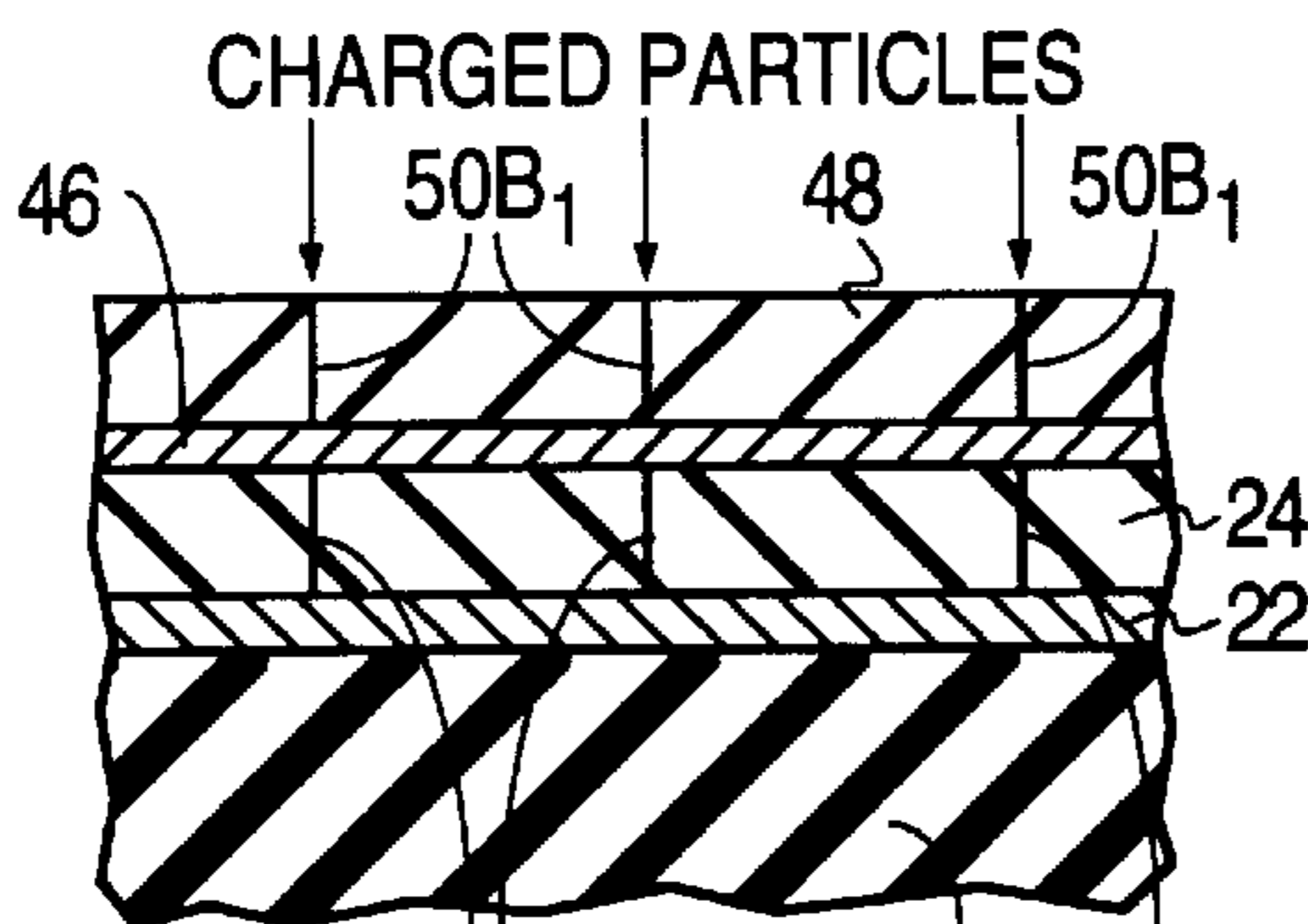


FIG. 5c

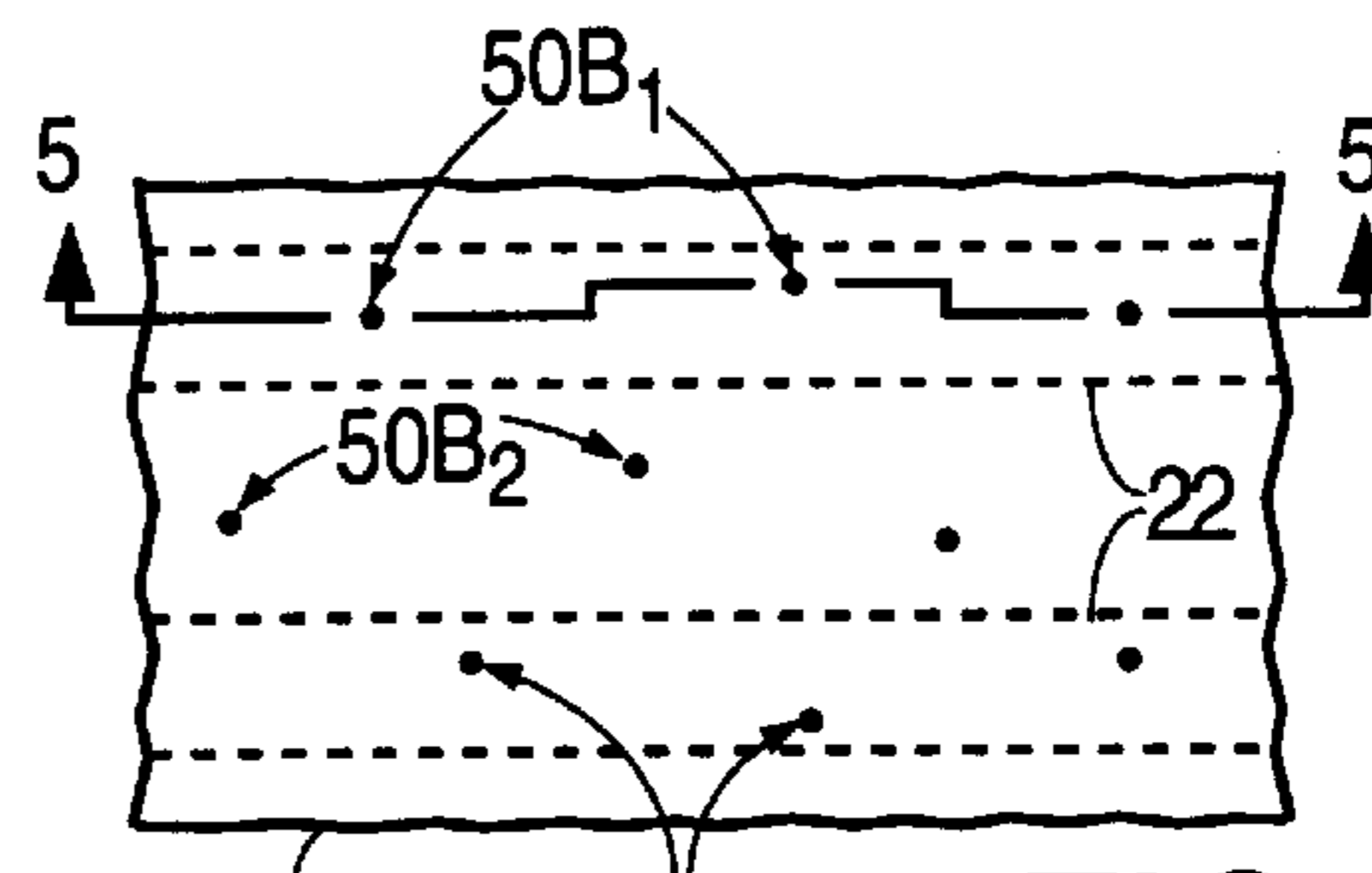


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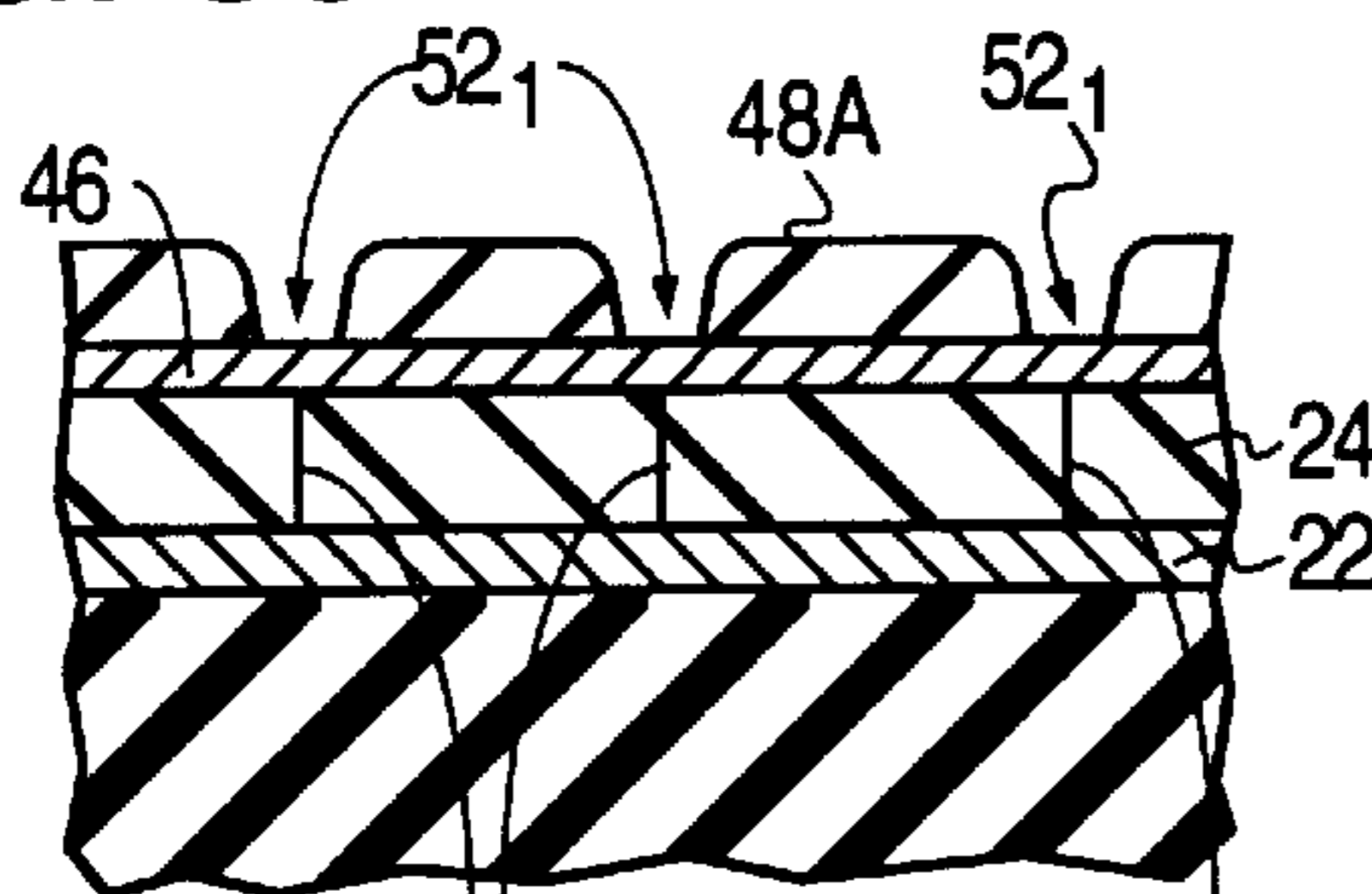


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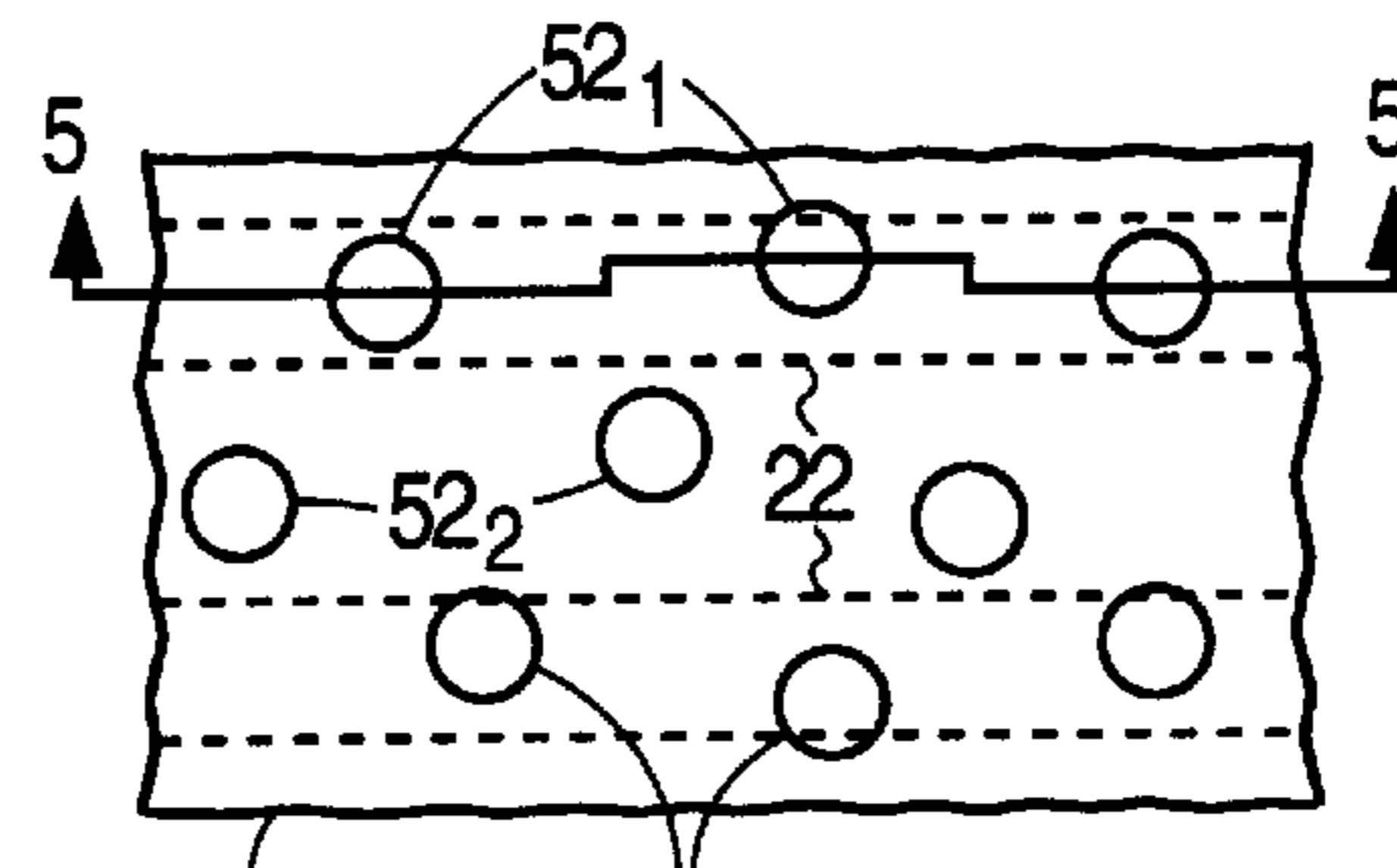


FIG. 6d

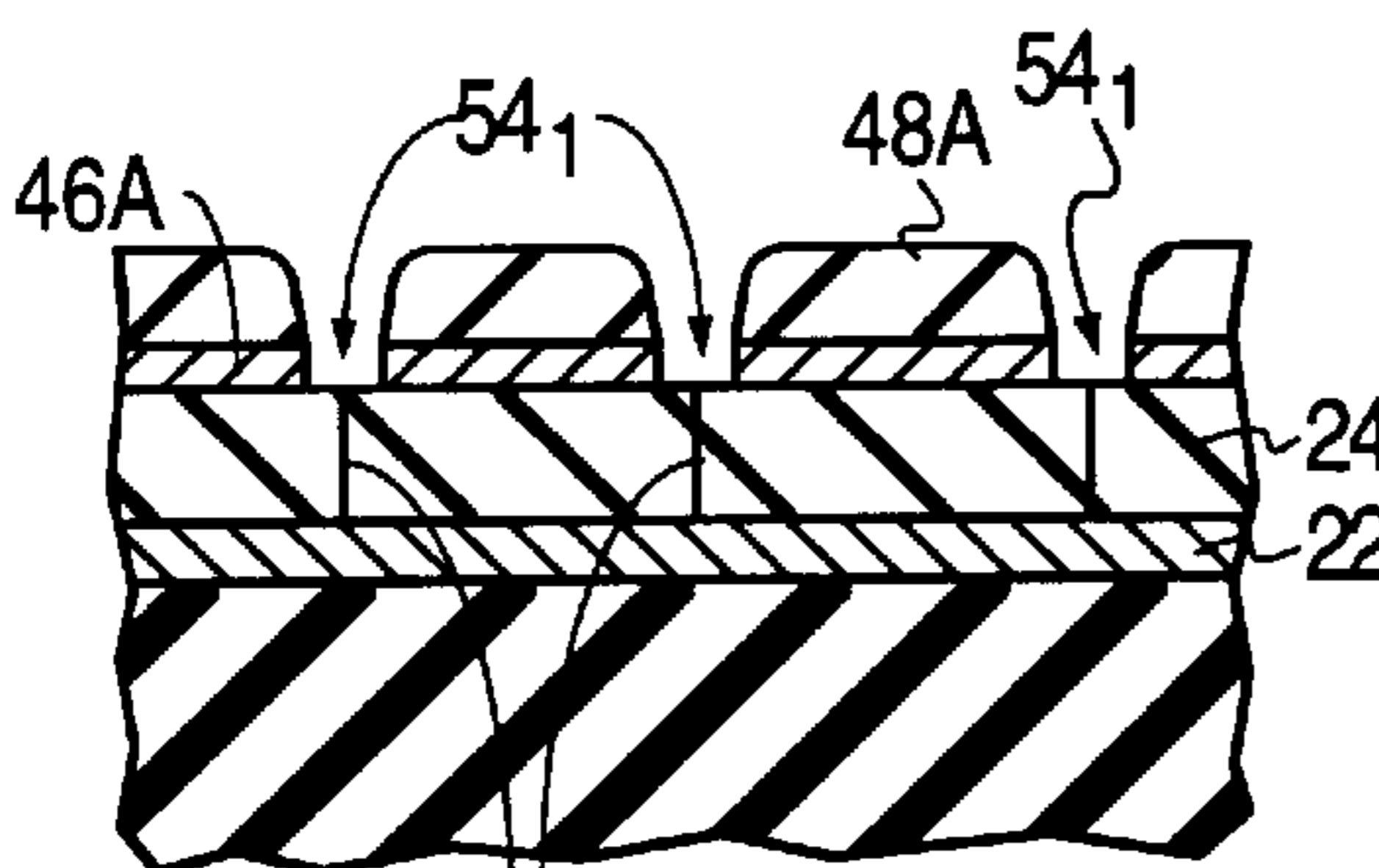


FIG. 5e

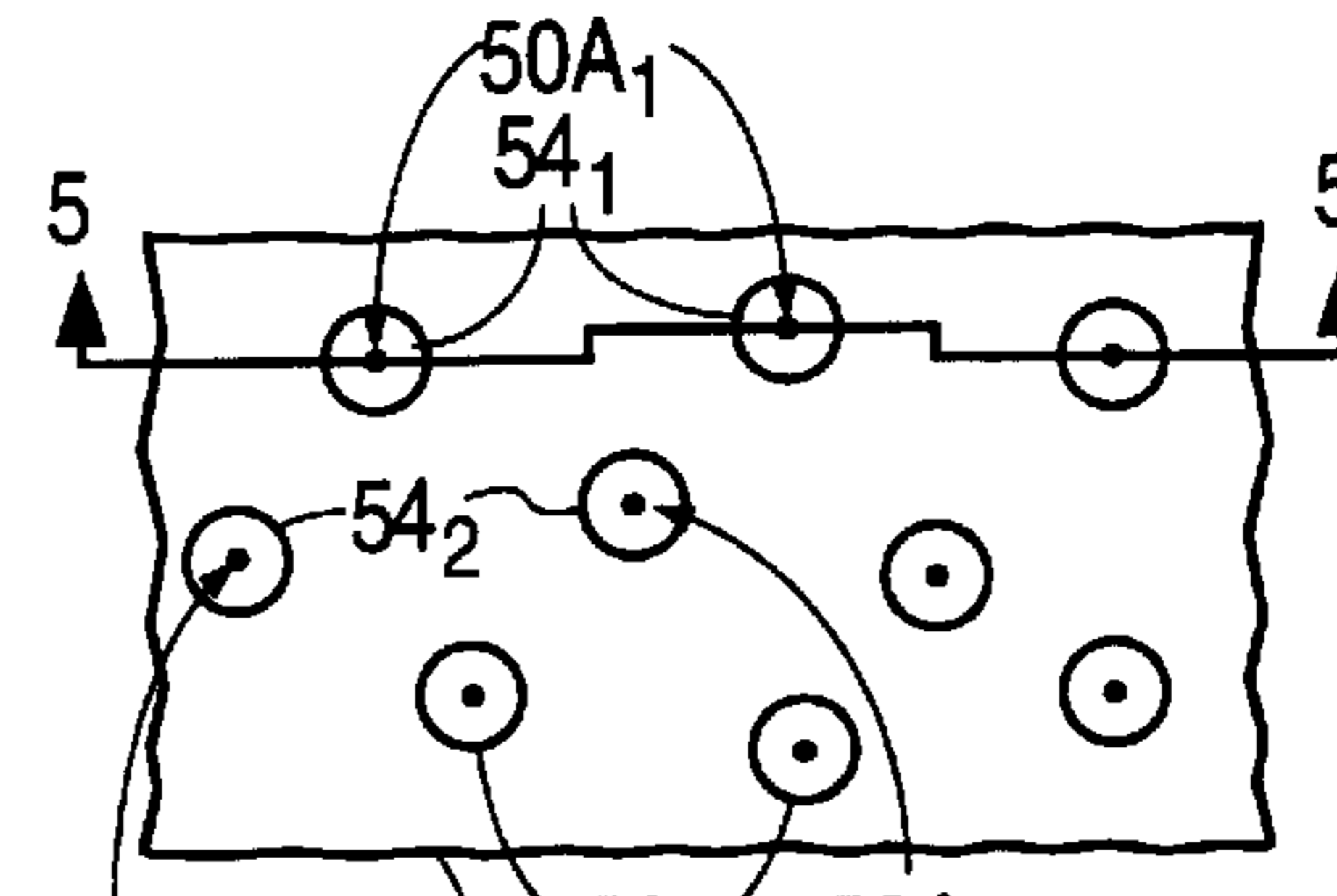


FIG. 6e

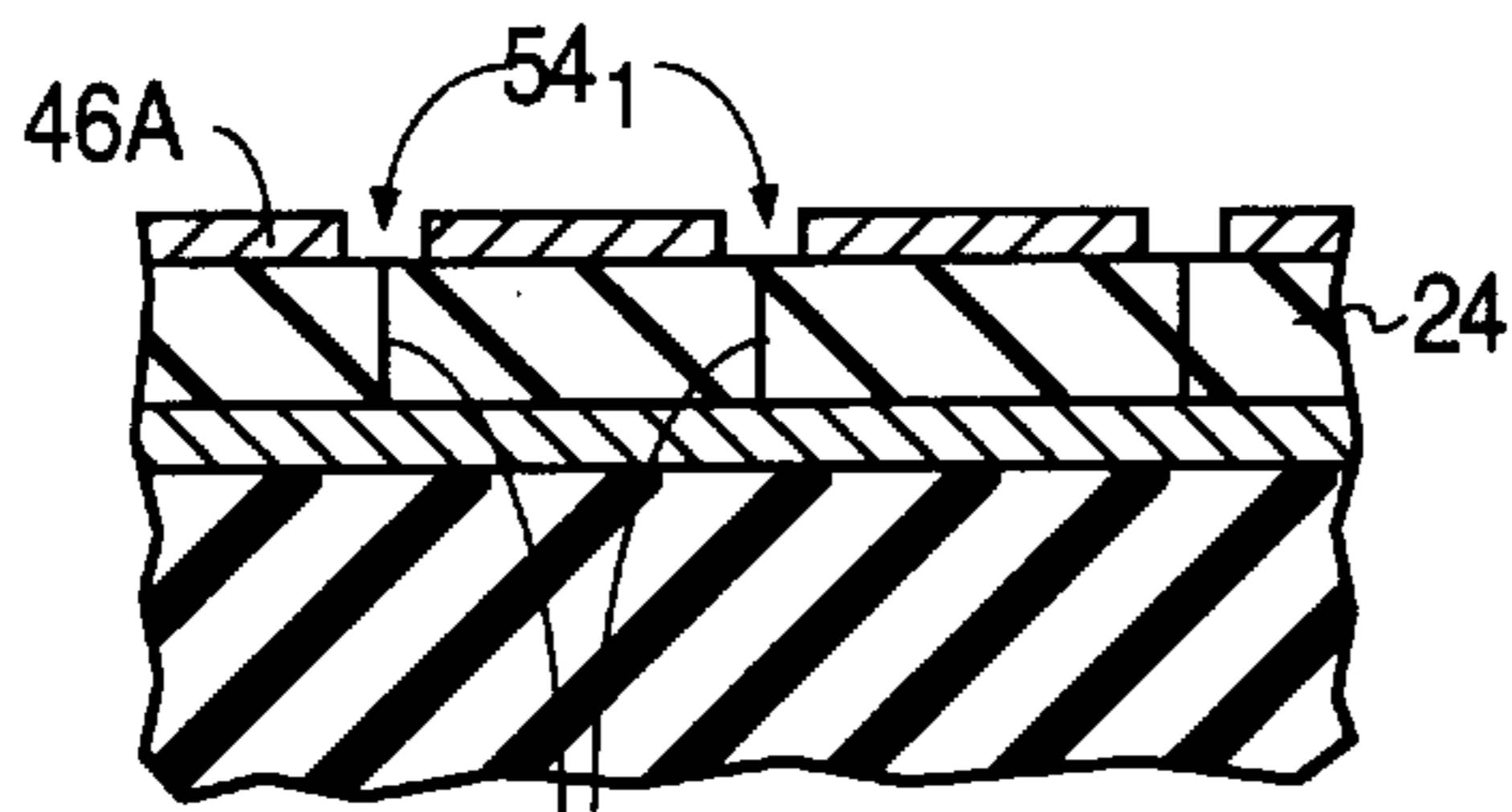


FIG. 5f

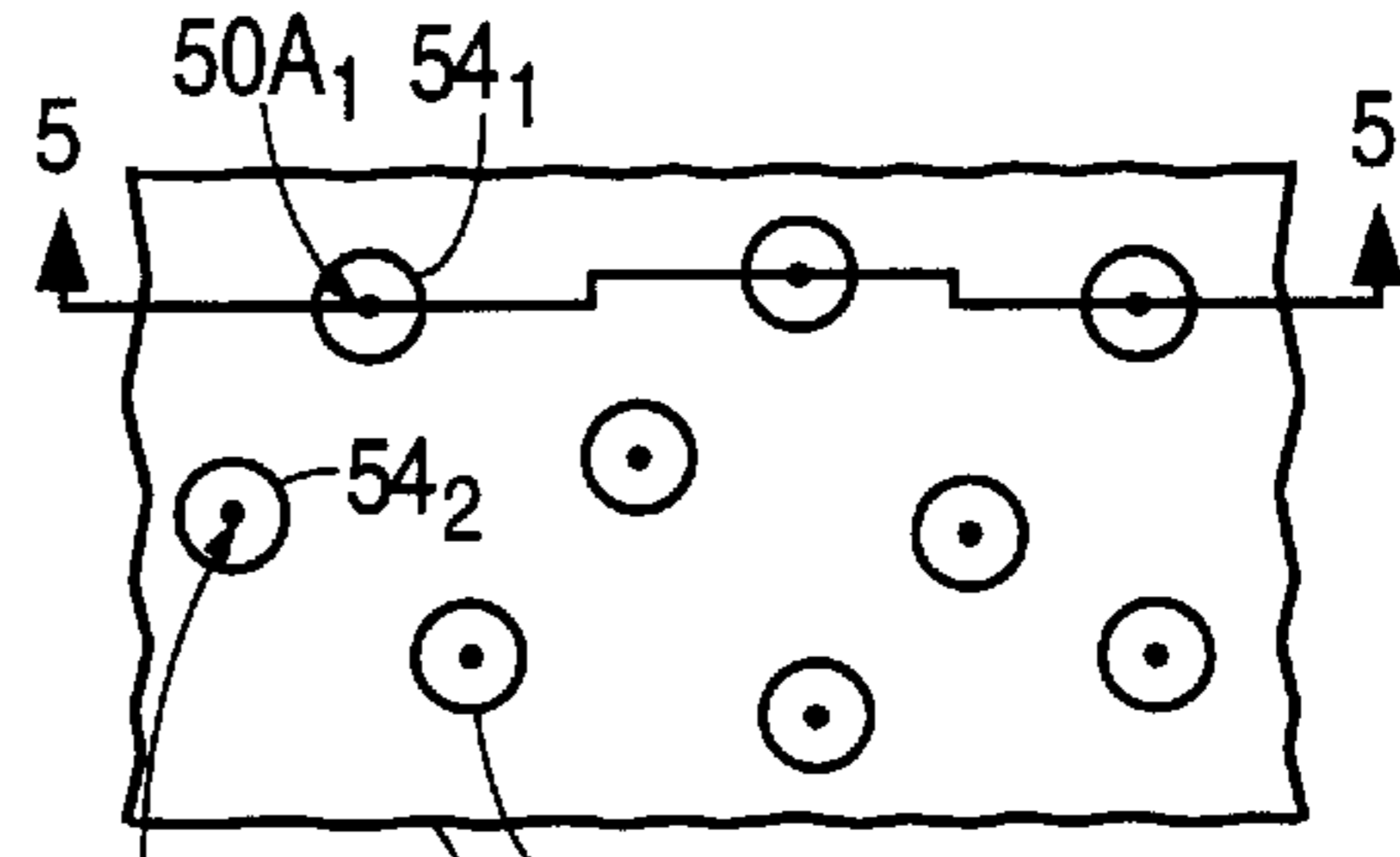


FIG. 6f

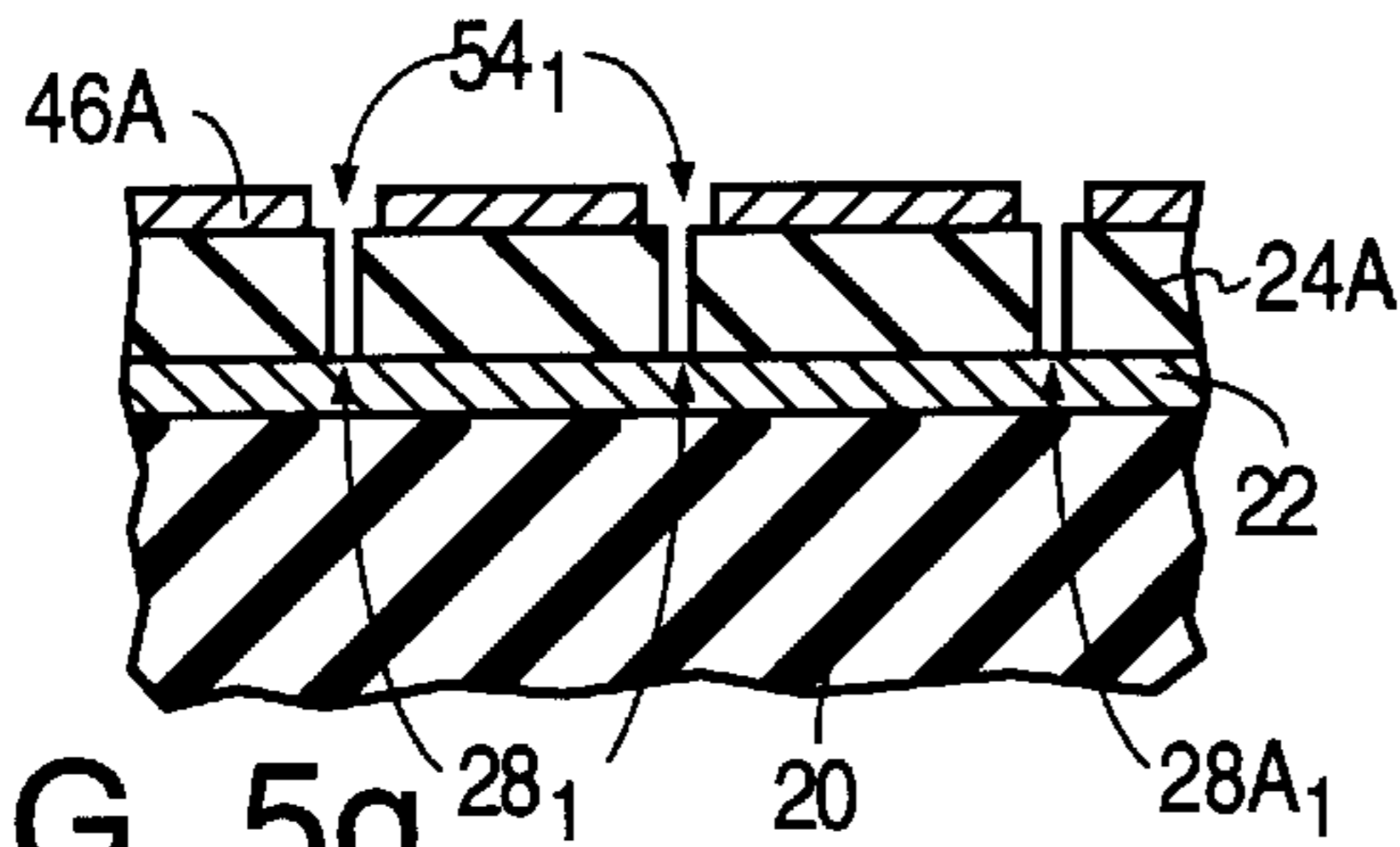


FIG. 5g

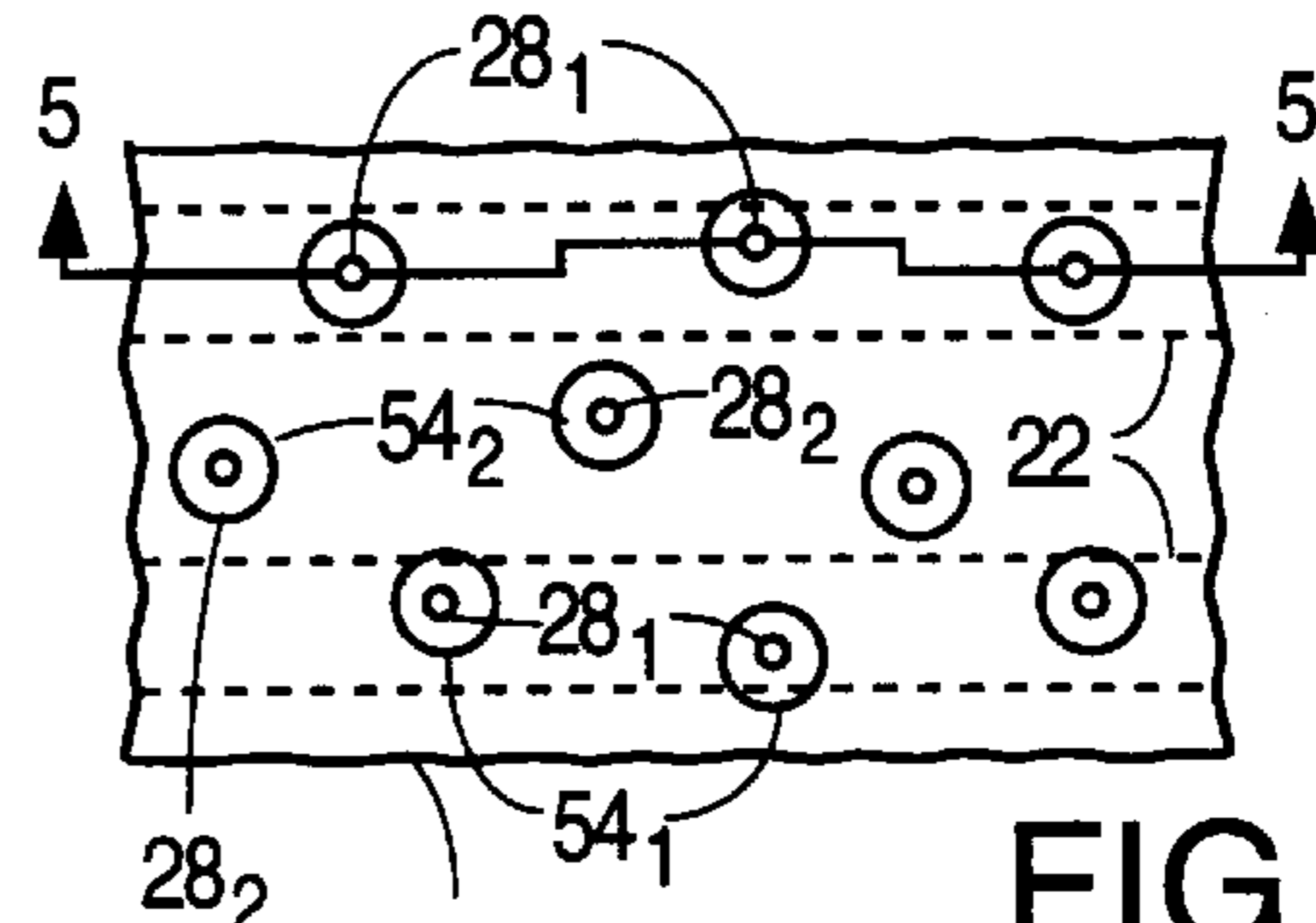


FIG. 6g

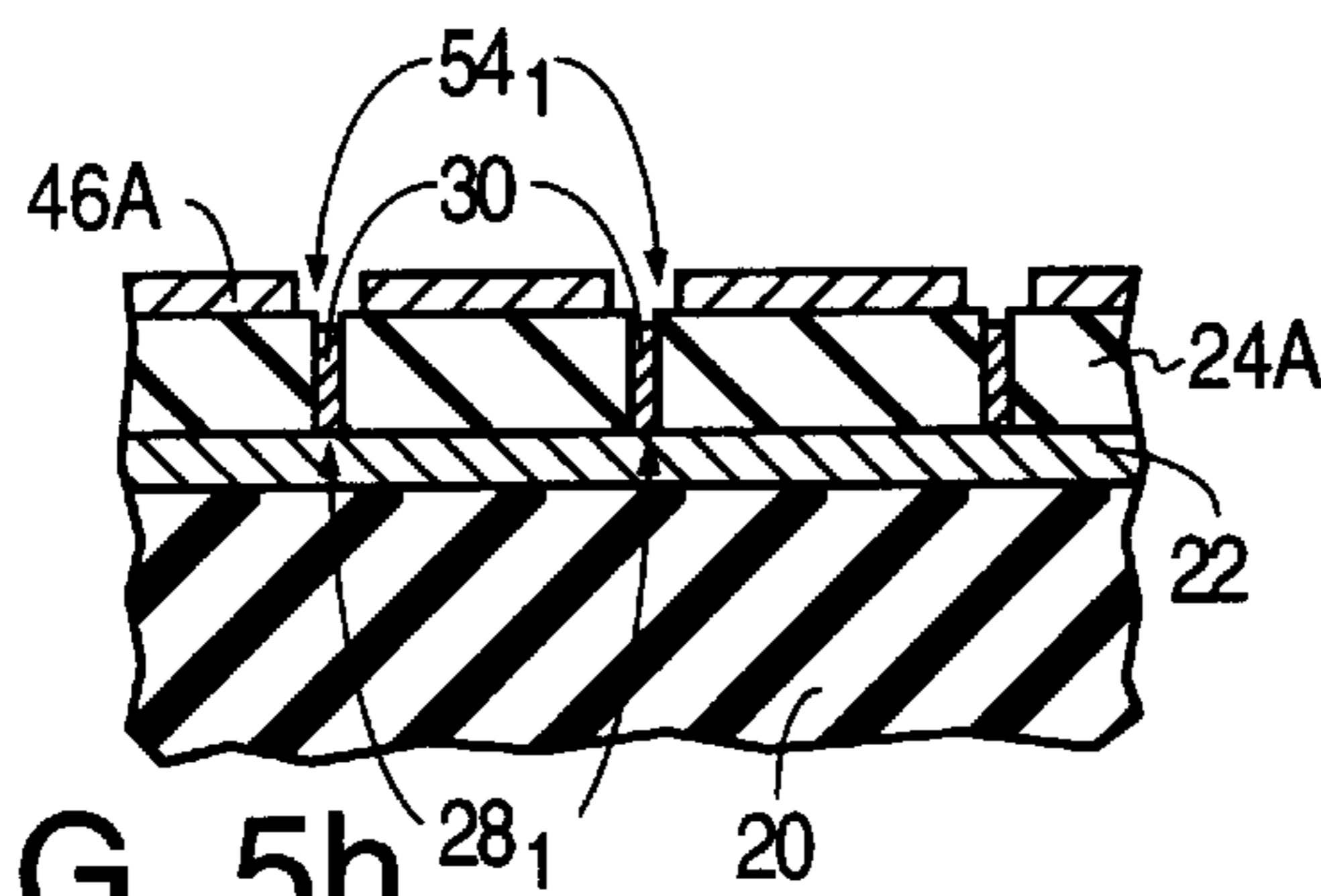


FIG. 5h

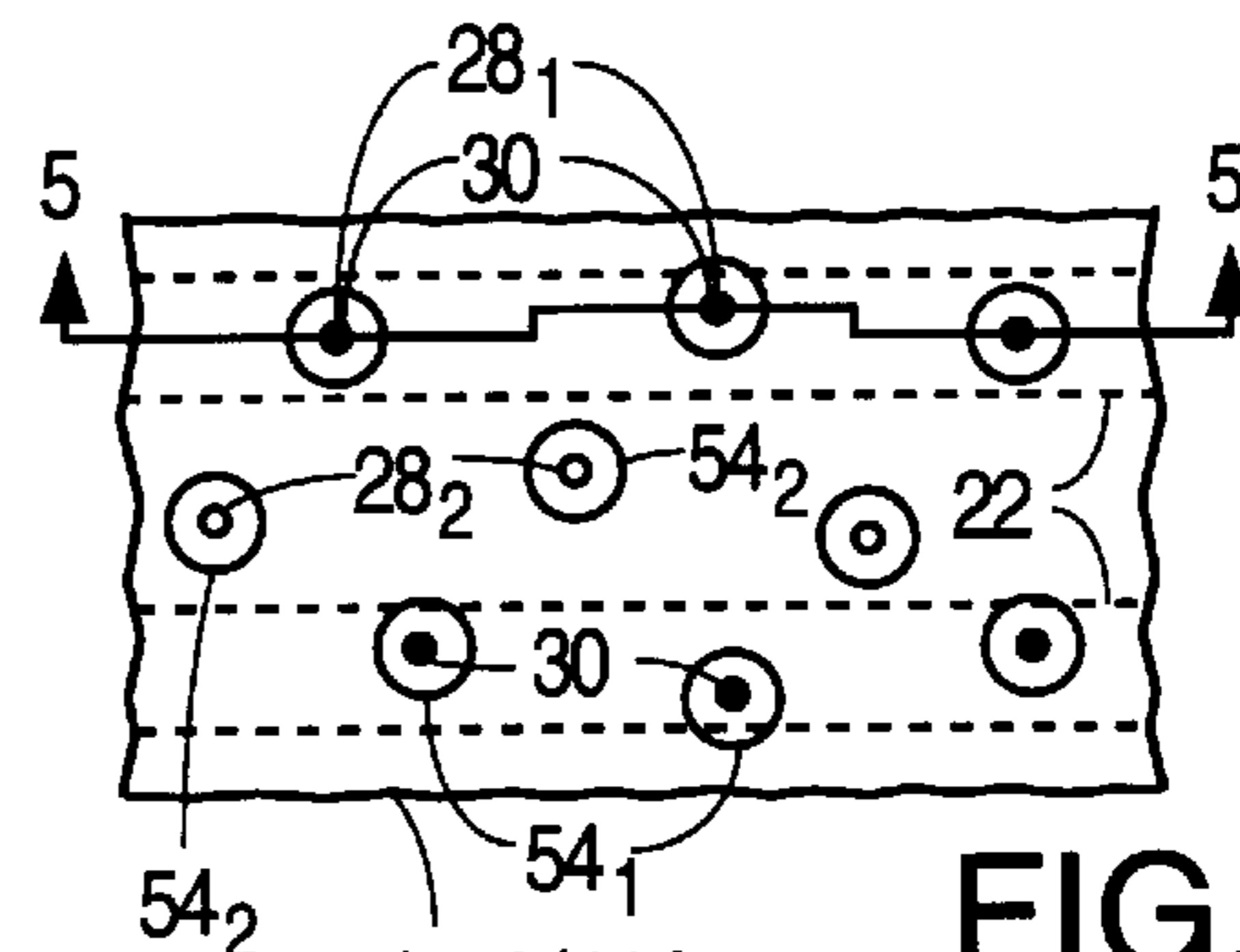


FIG. 6h

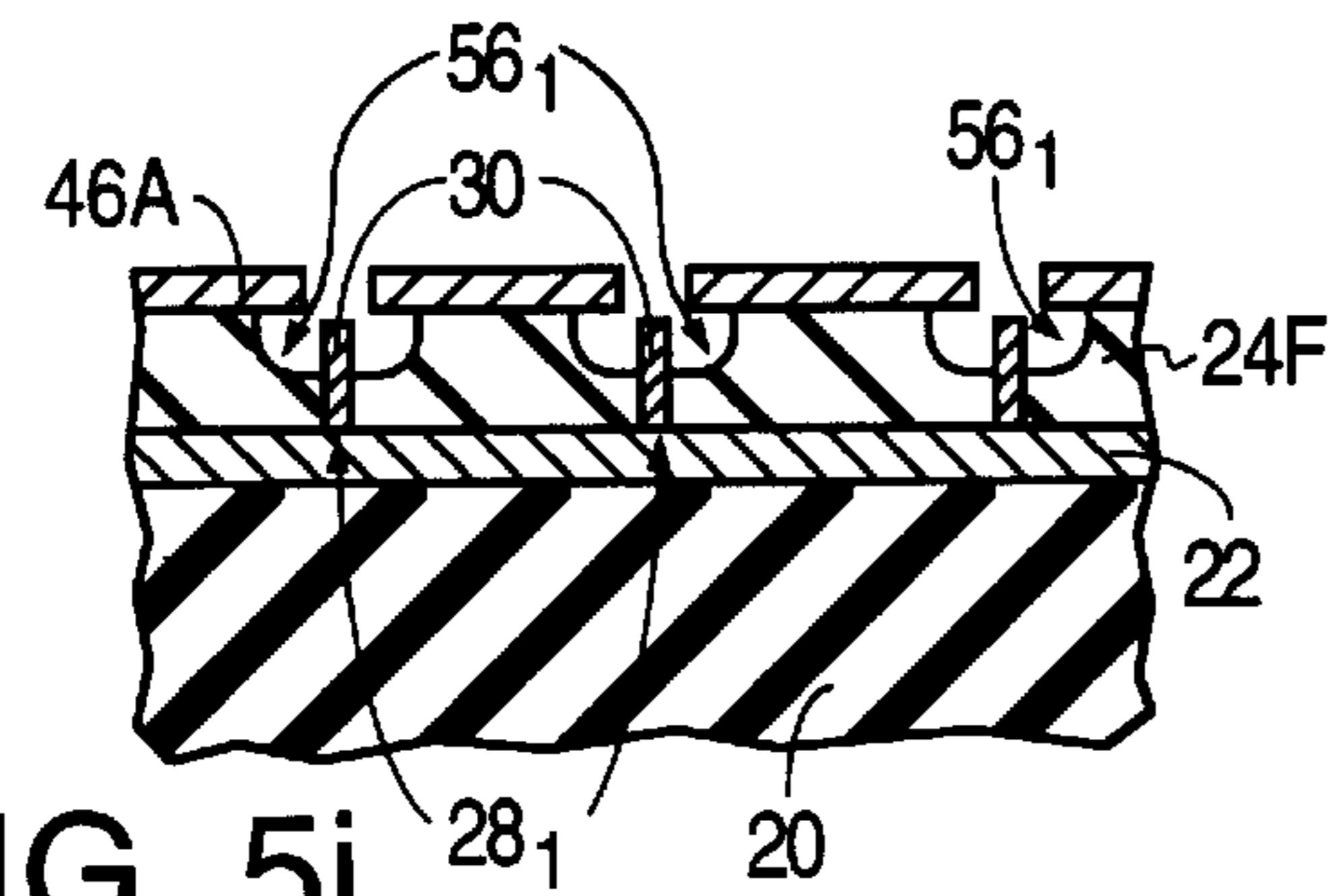


FIG. 5i

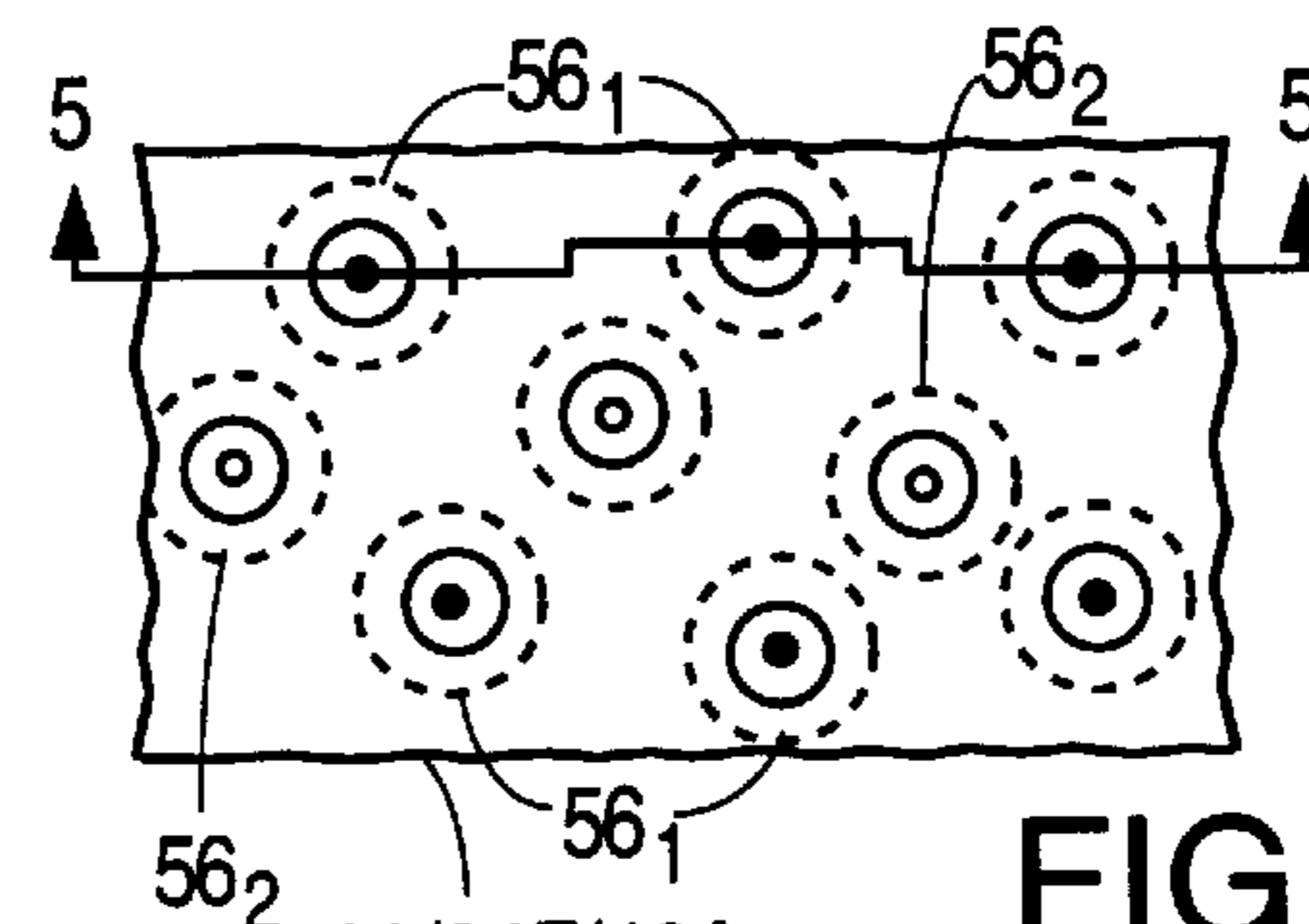


FIG. 6i

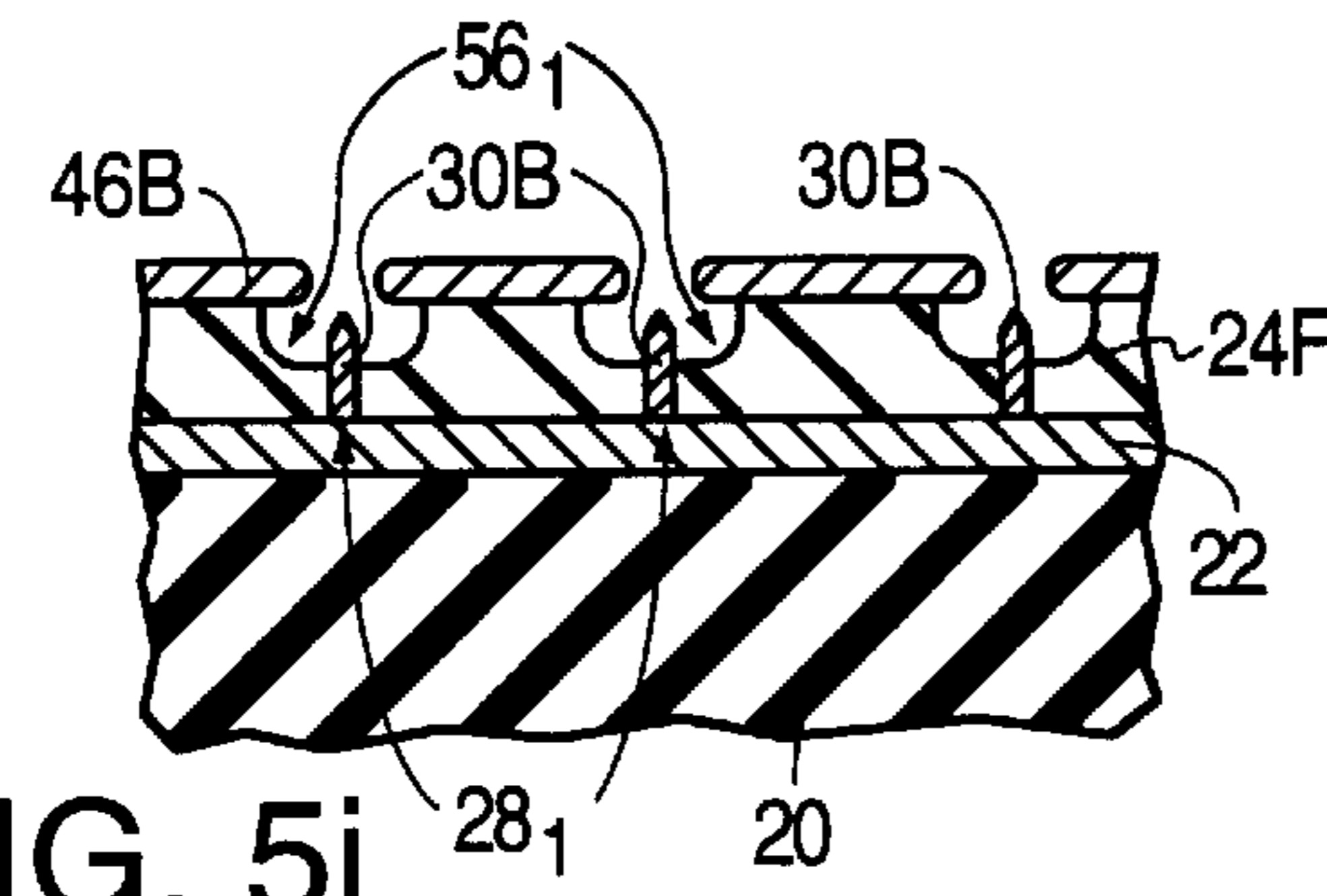


FIG. 5j

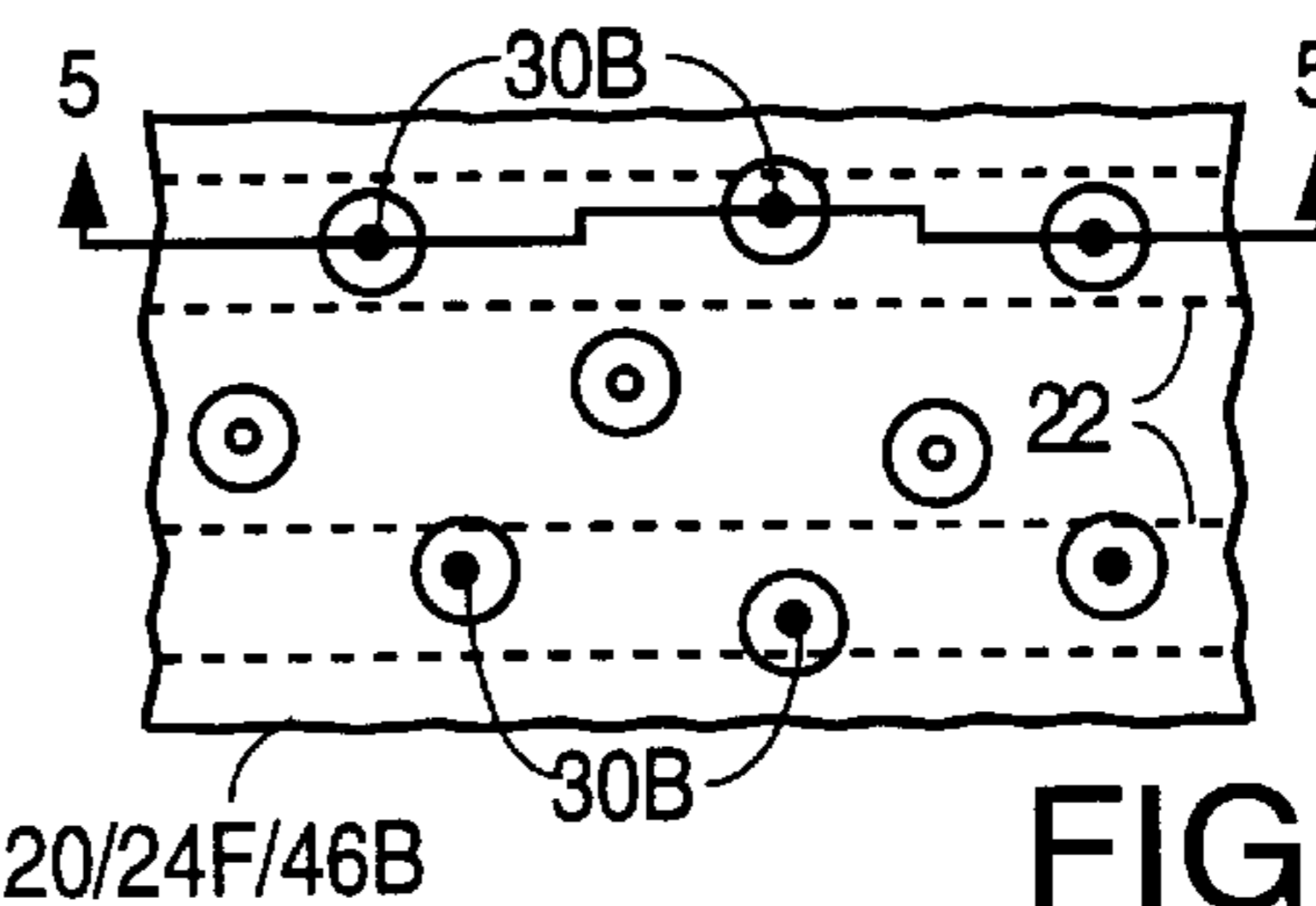
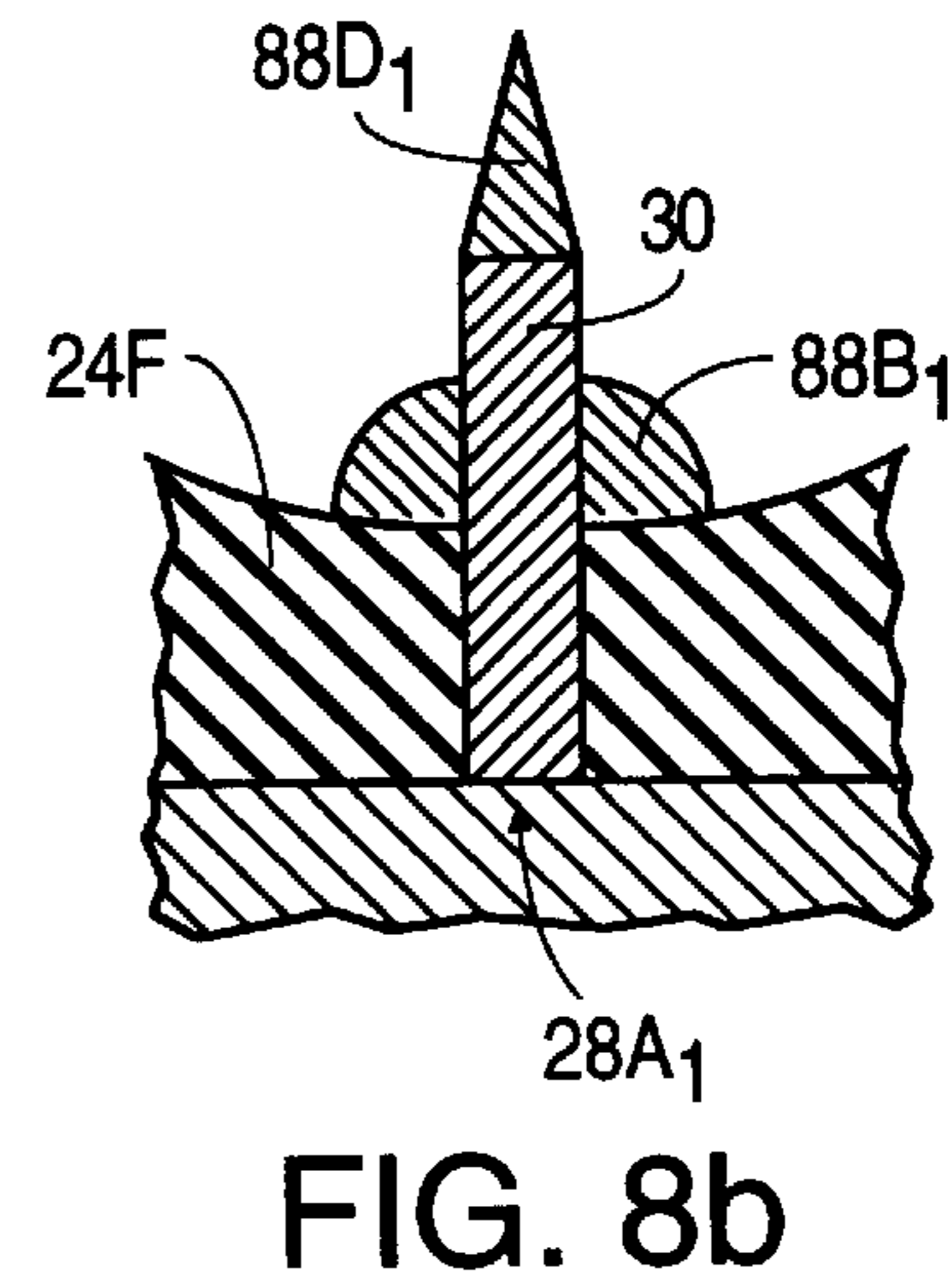
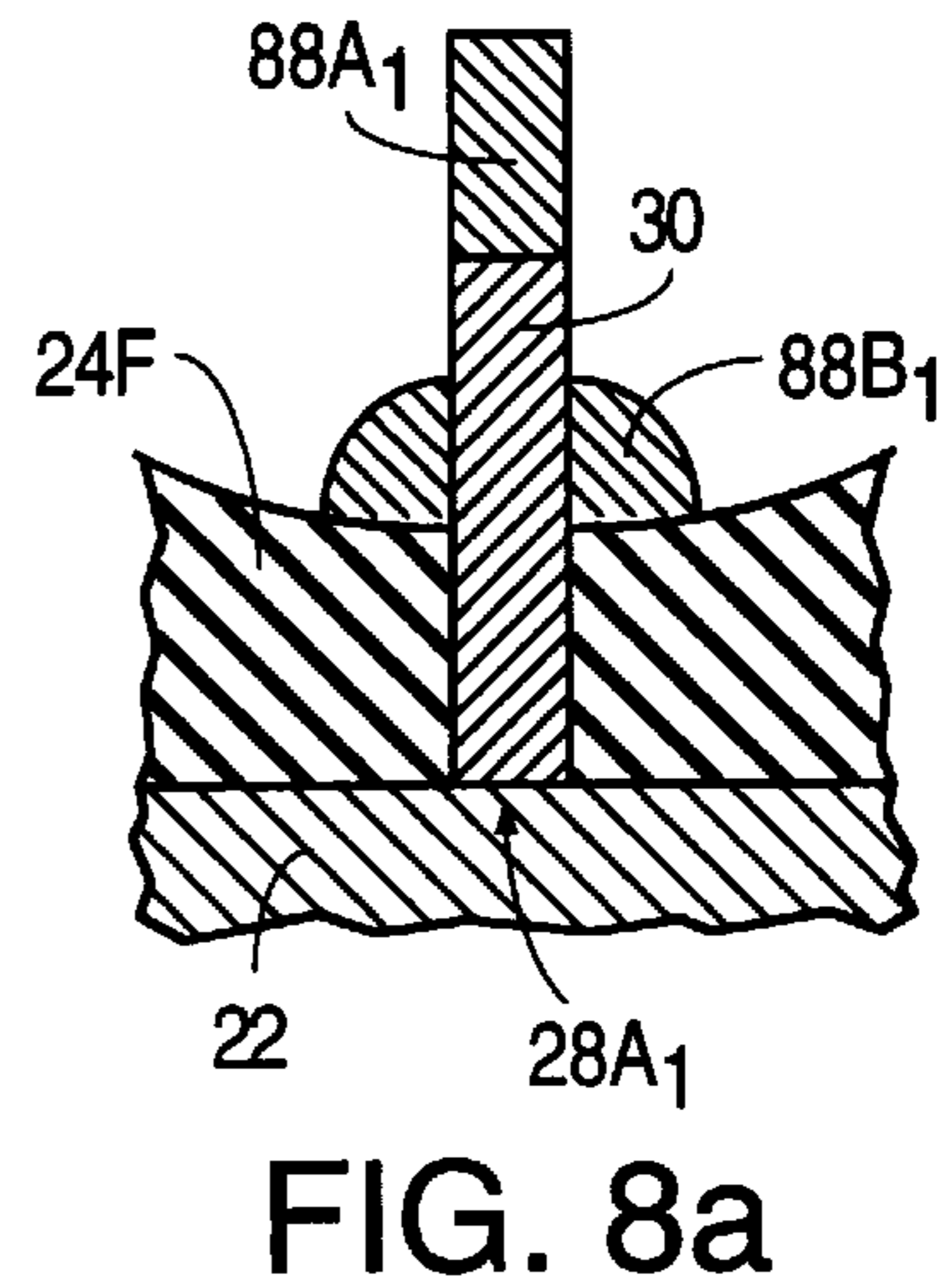
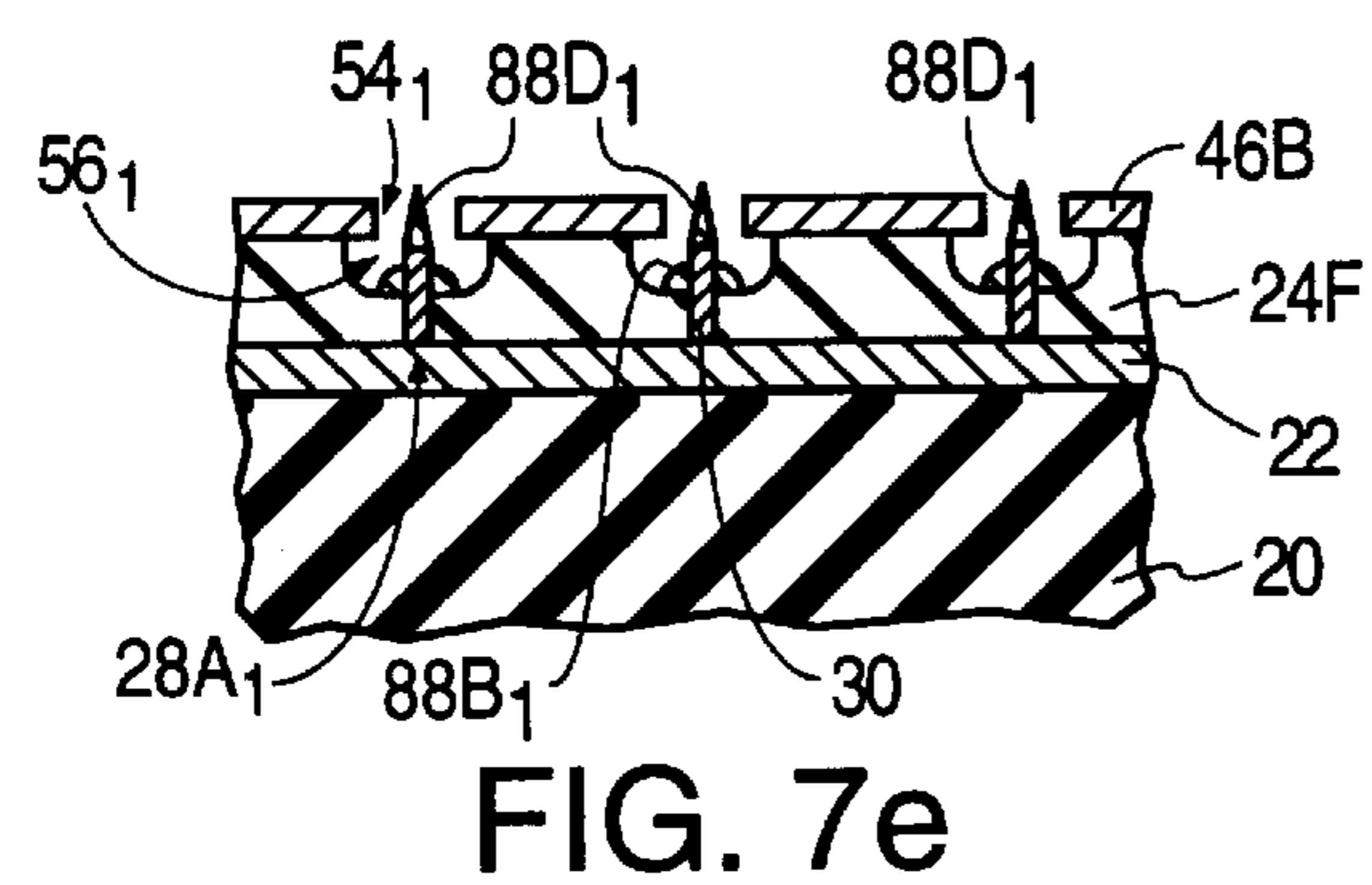
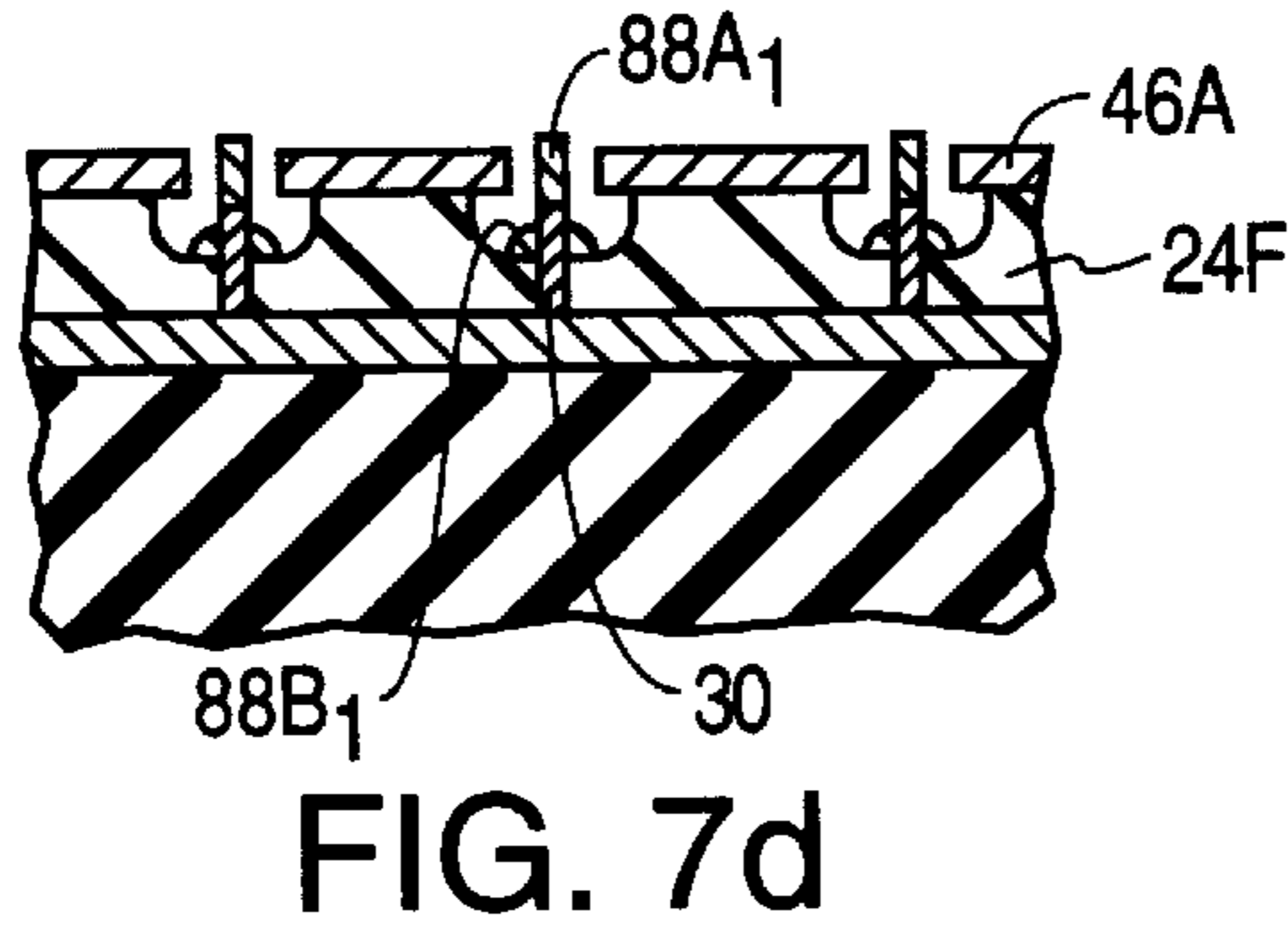
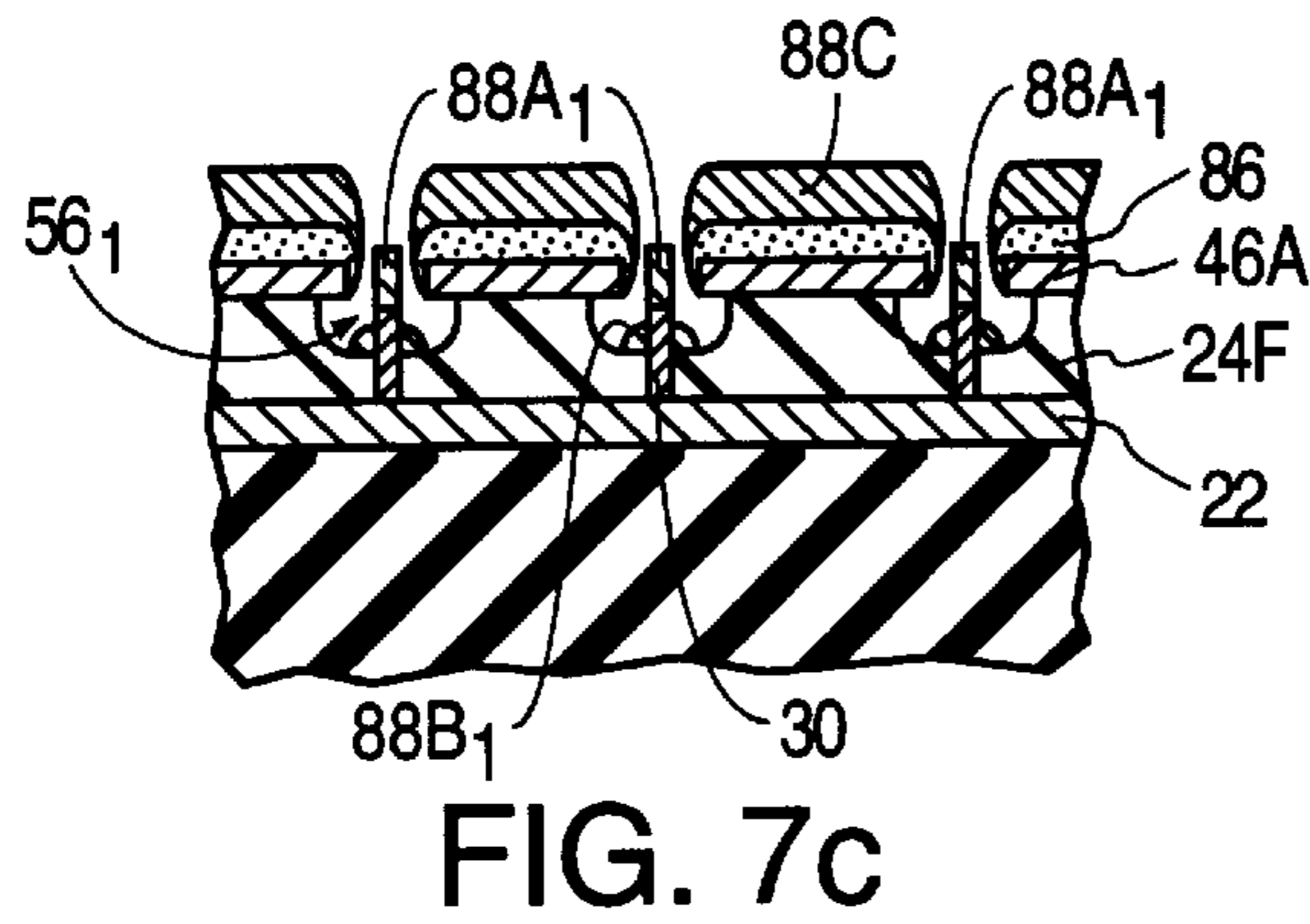
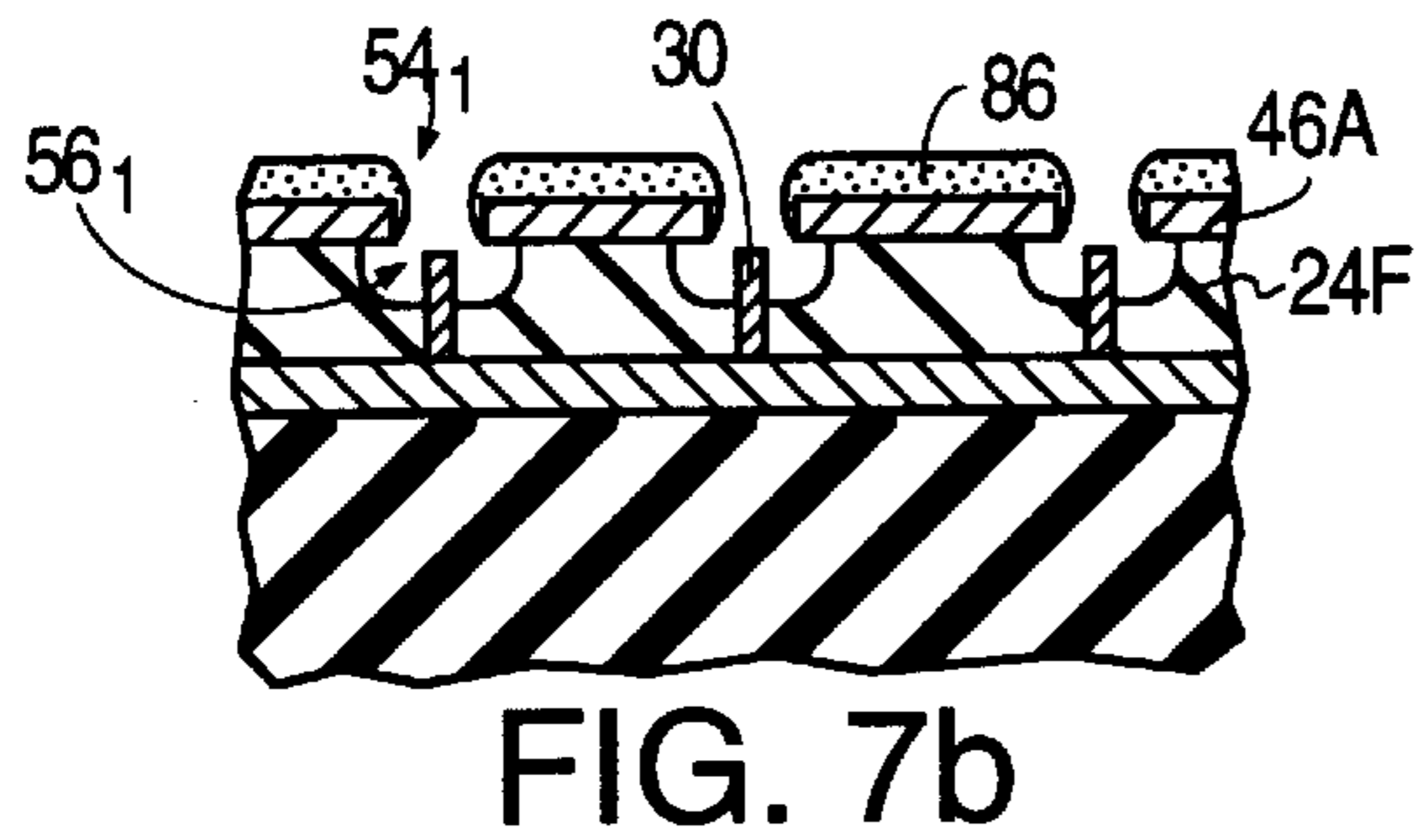
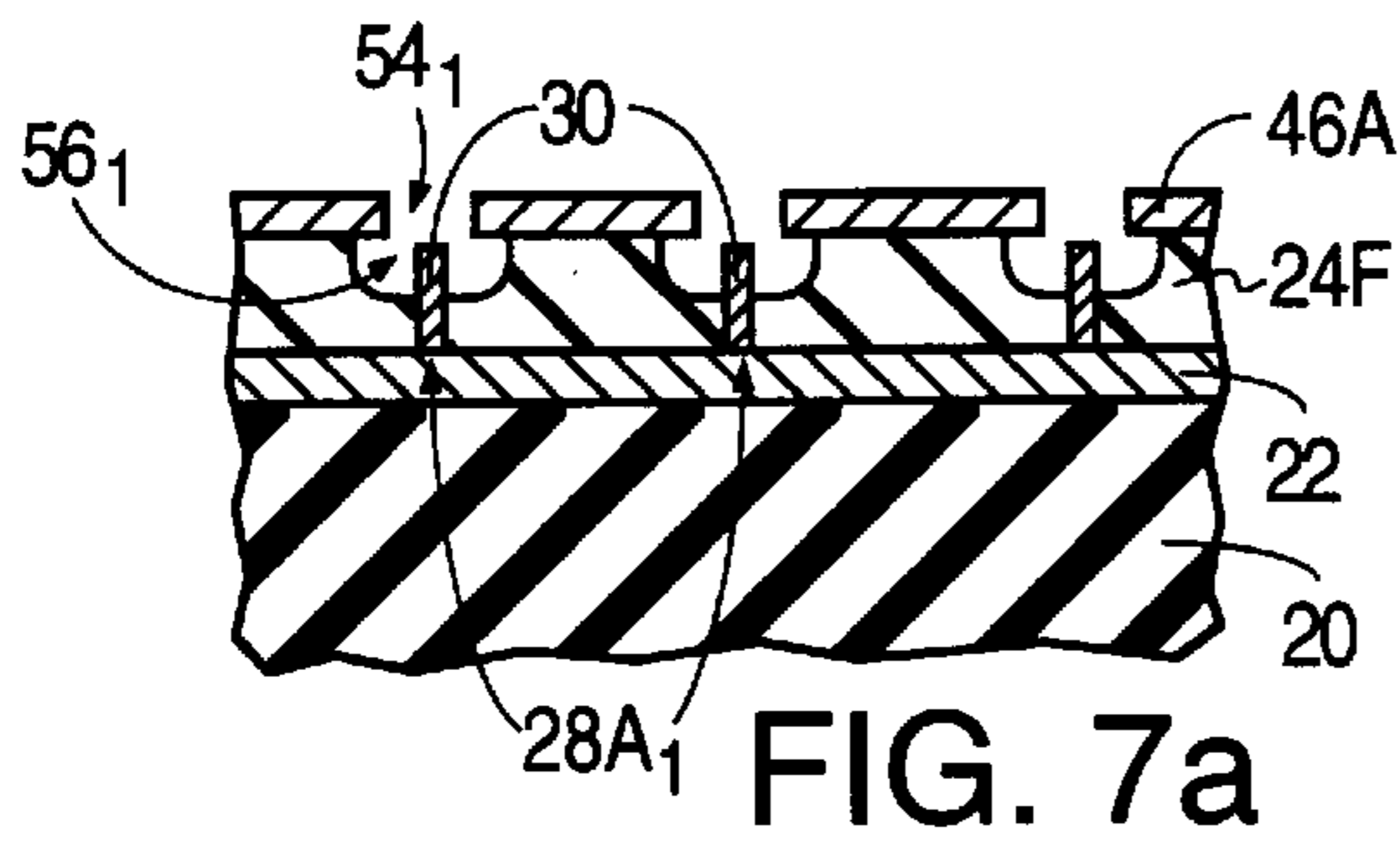


FIG. 6j



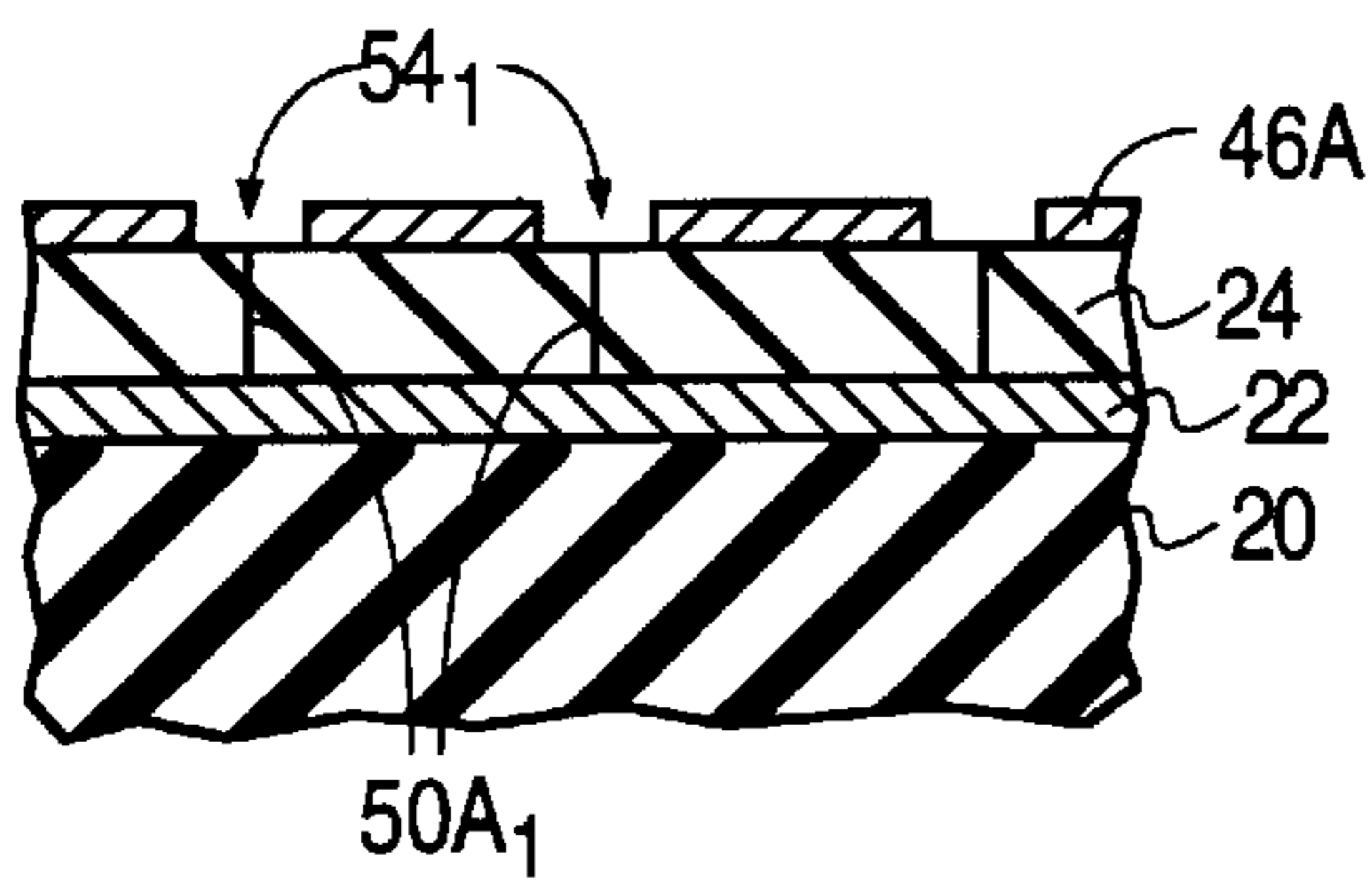


FIG. 9a

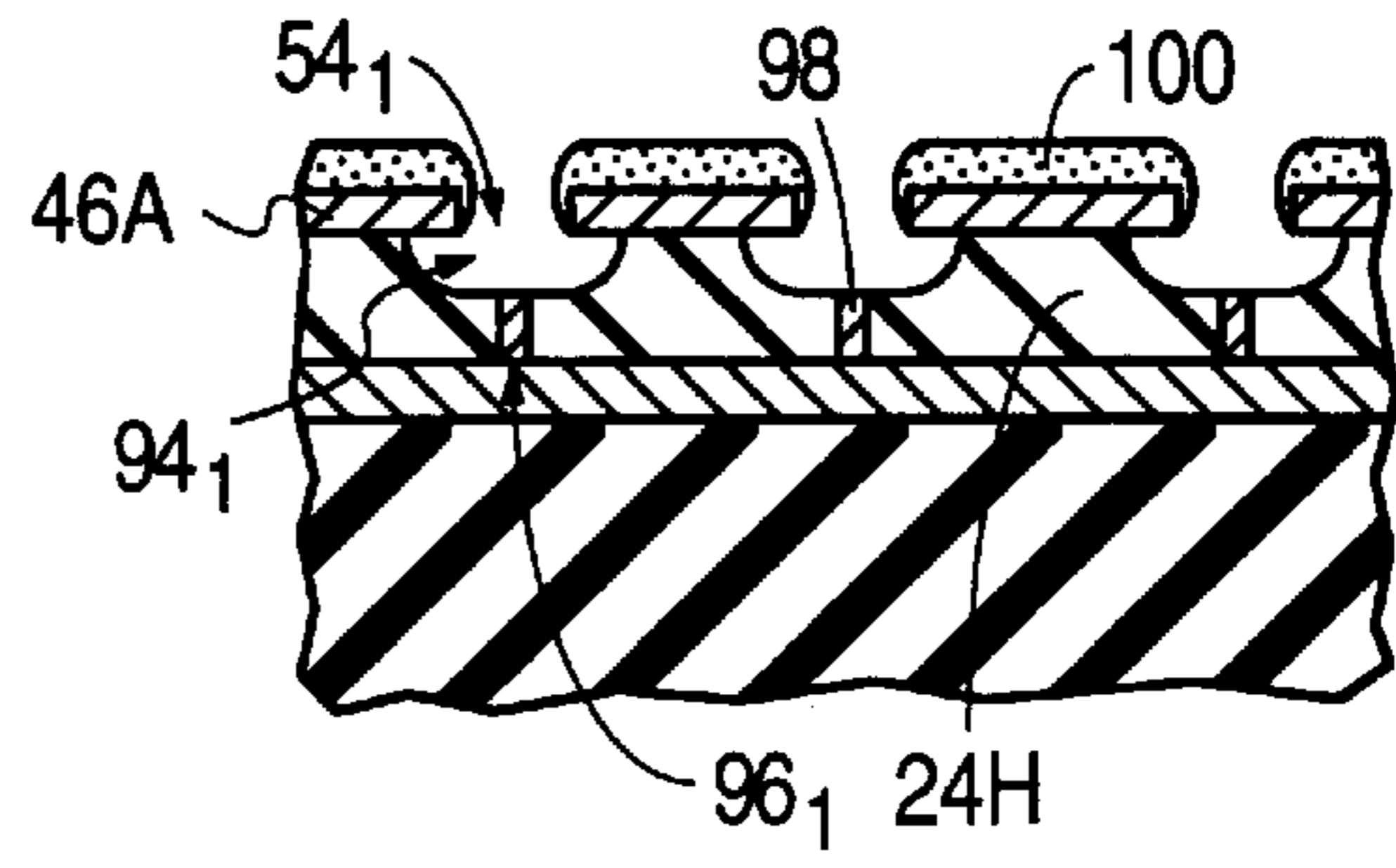


FIG. 9e

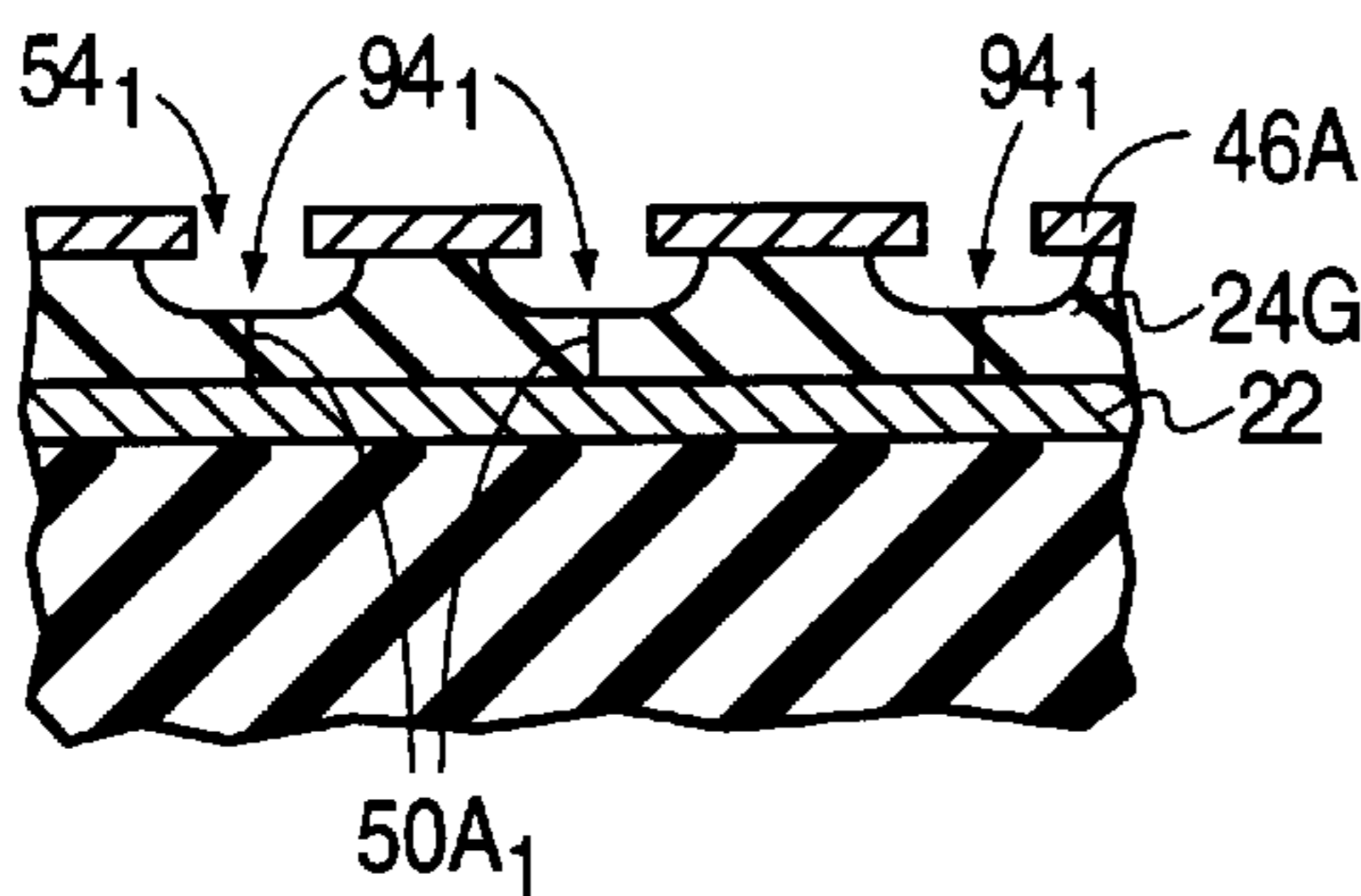


FIG. 9b

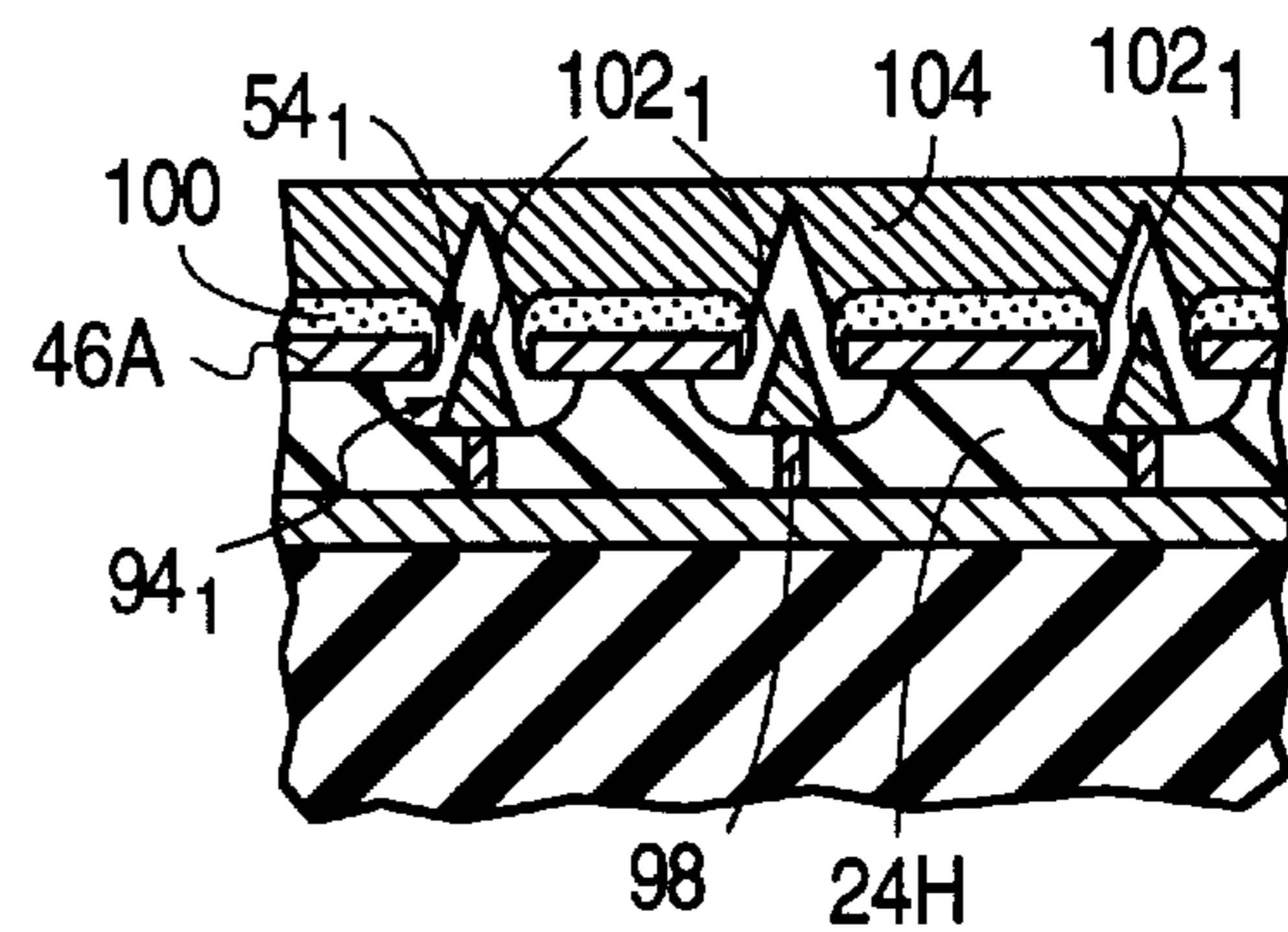


FIG. 9f

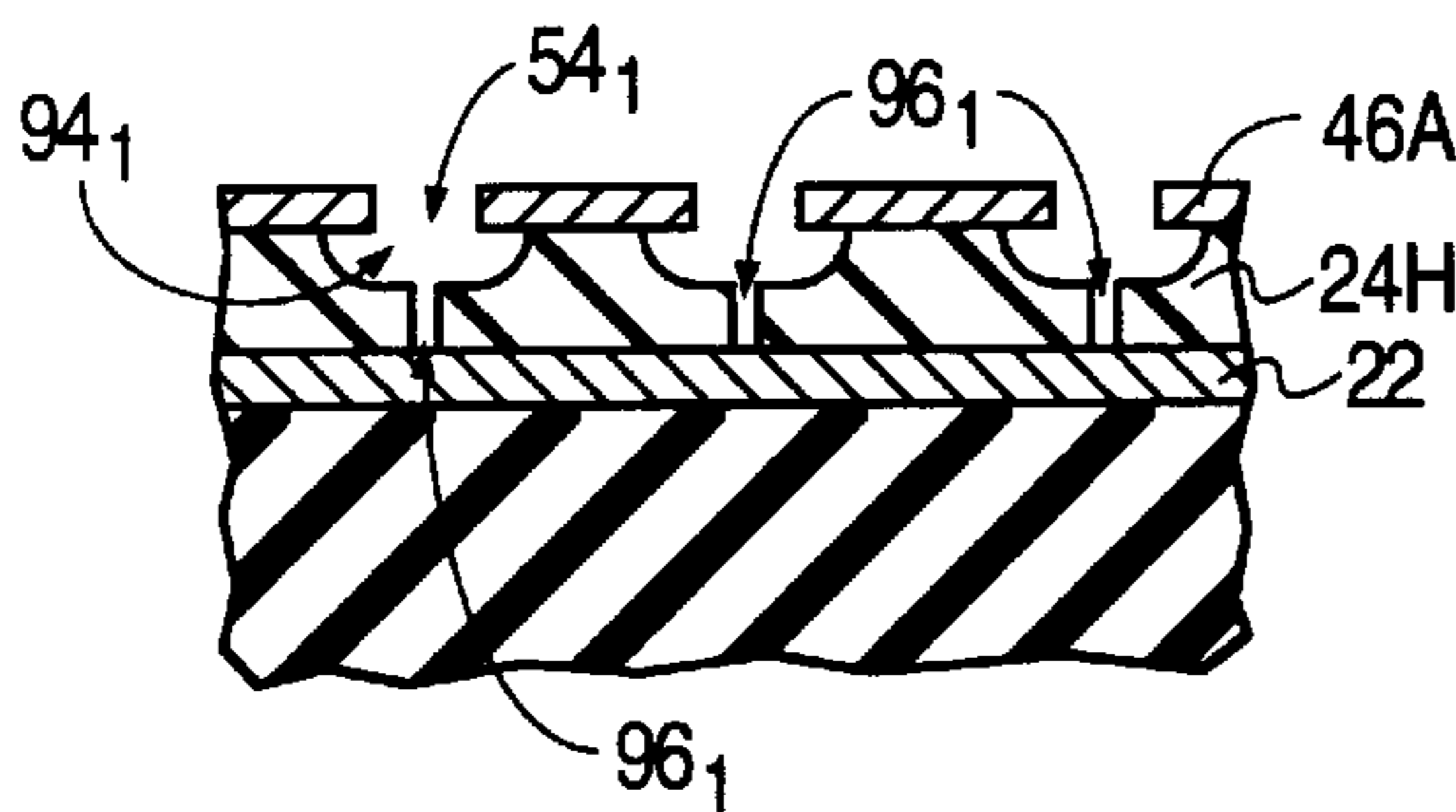


FIG. 9c

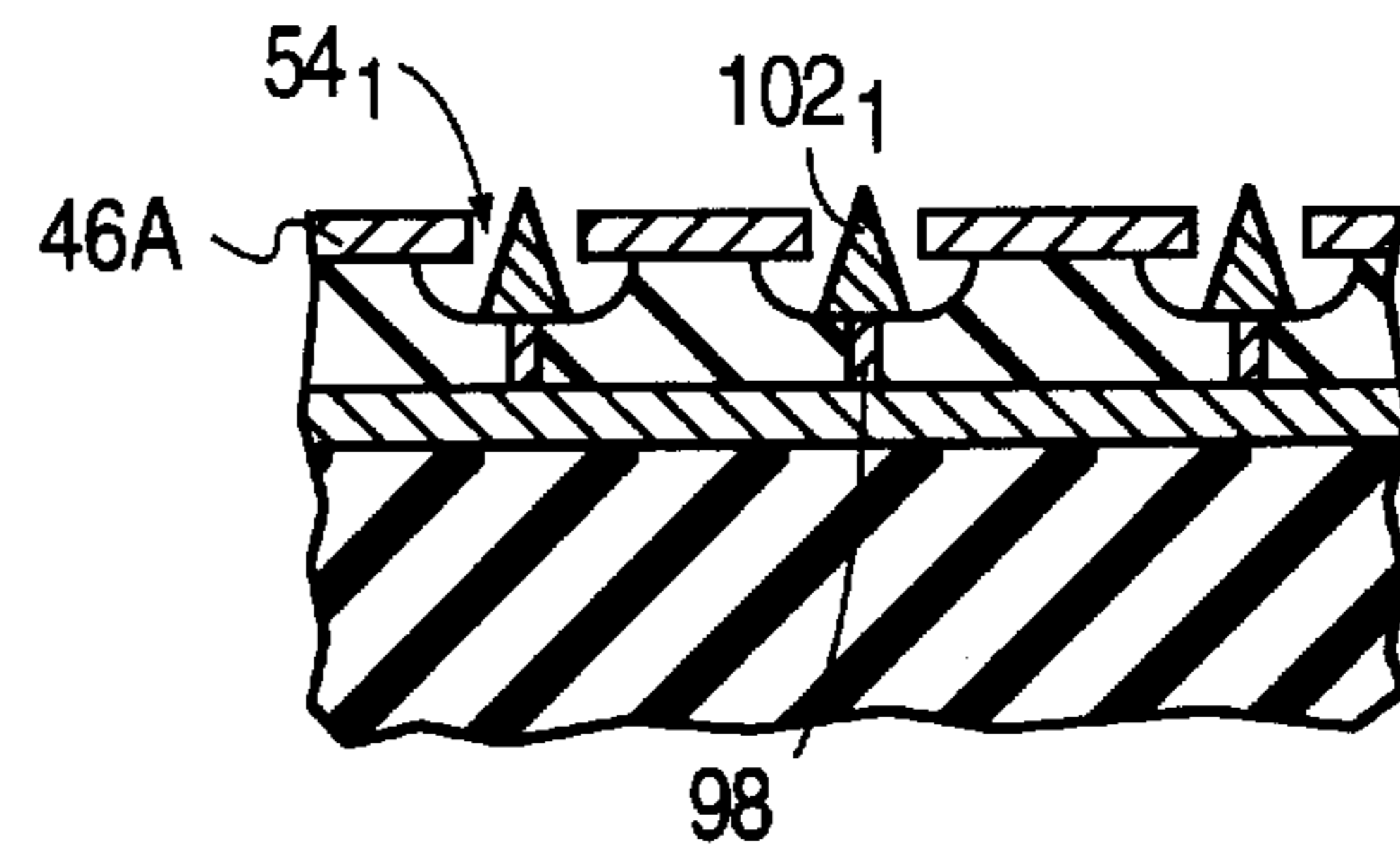


FIG. 9g

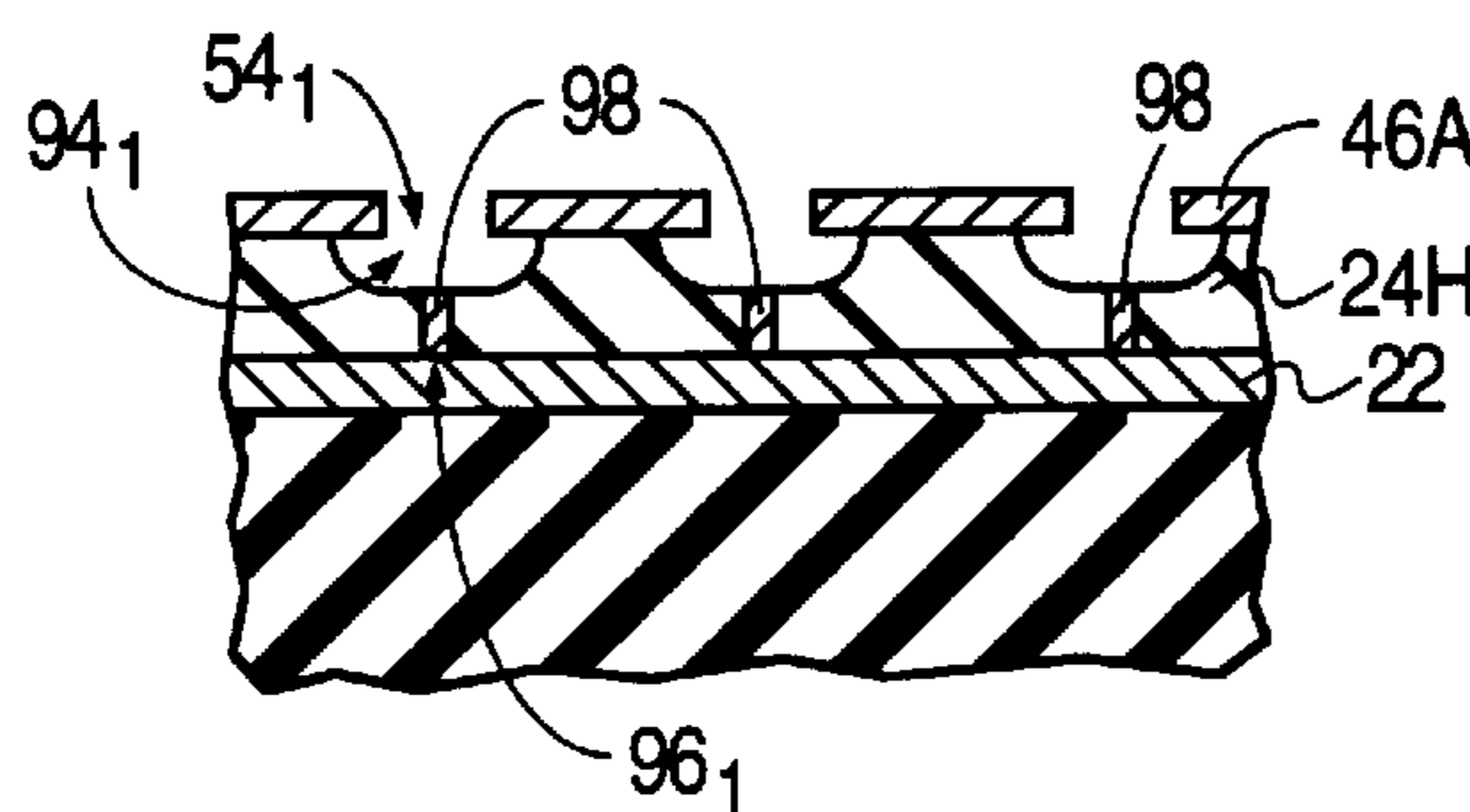


FIG. 9d

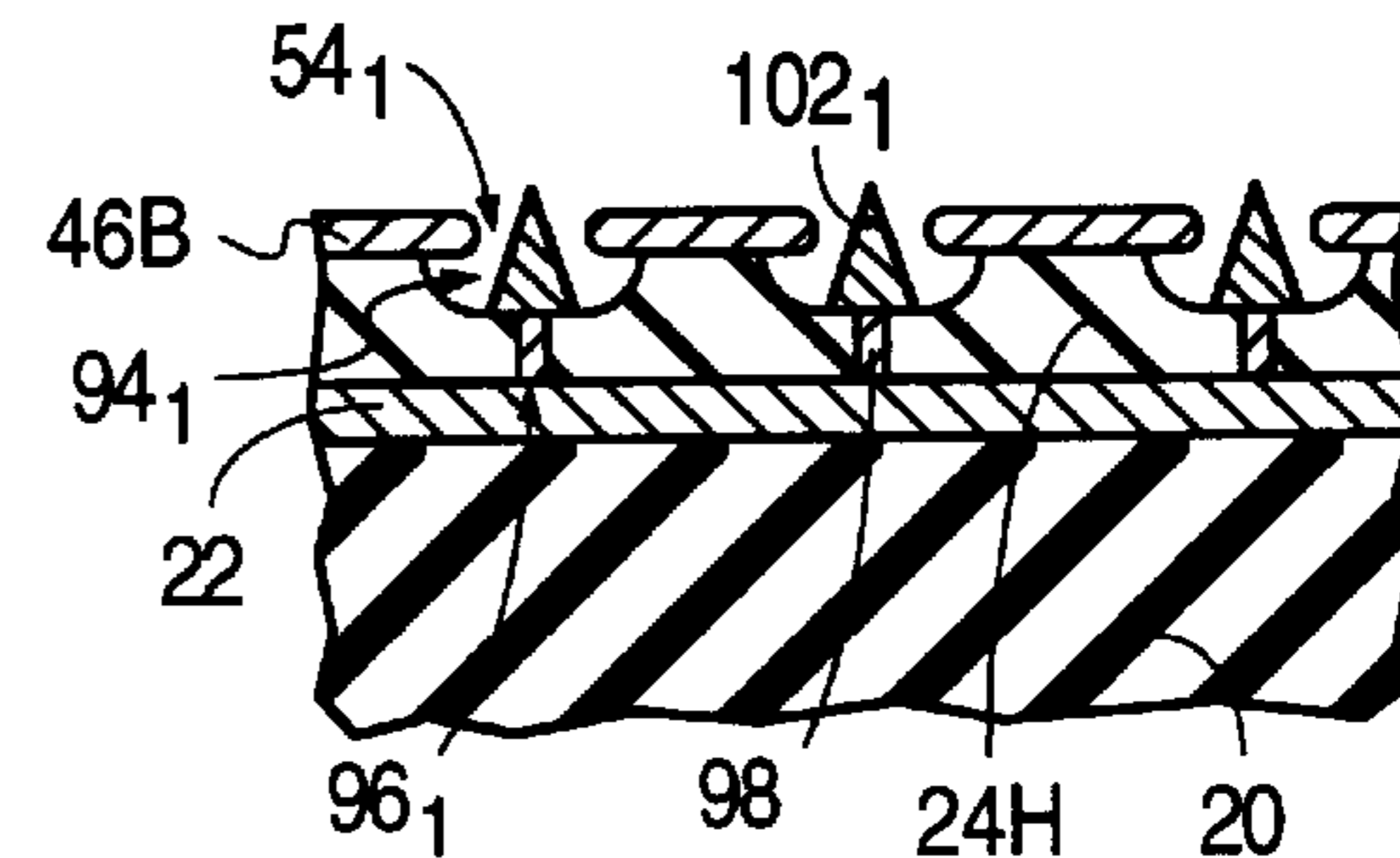
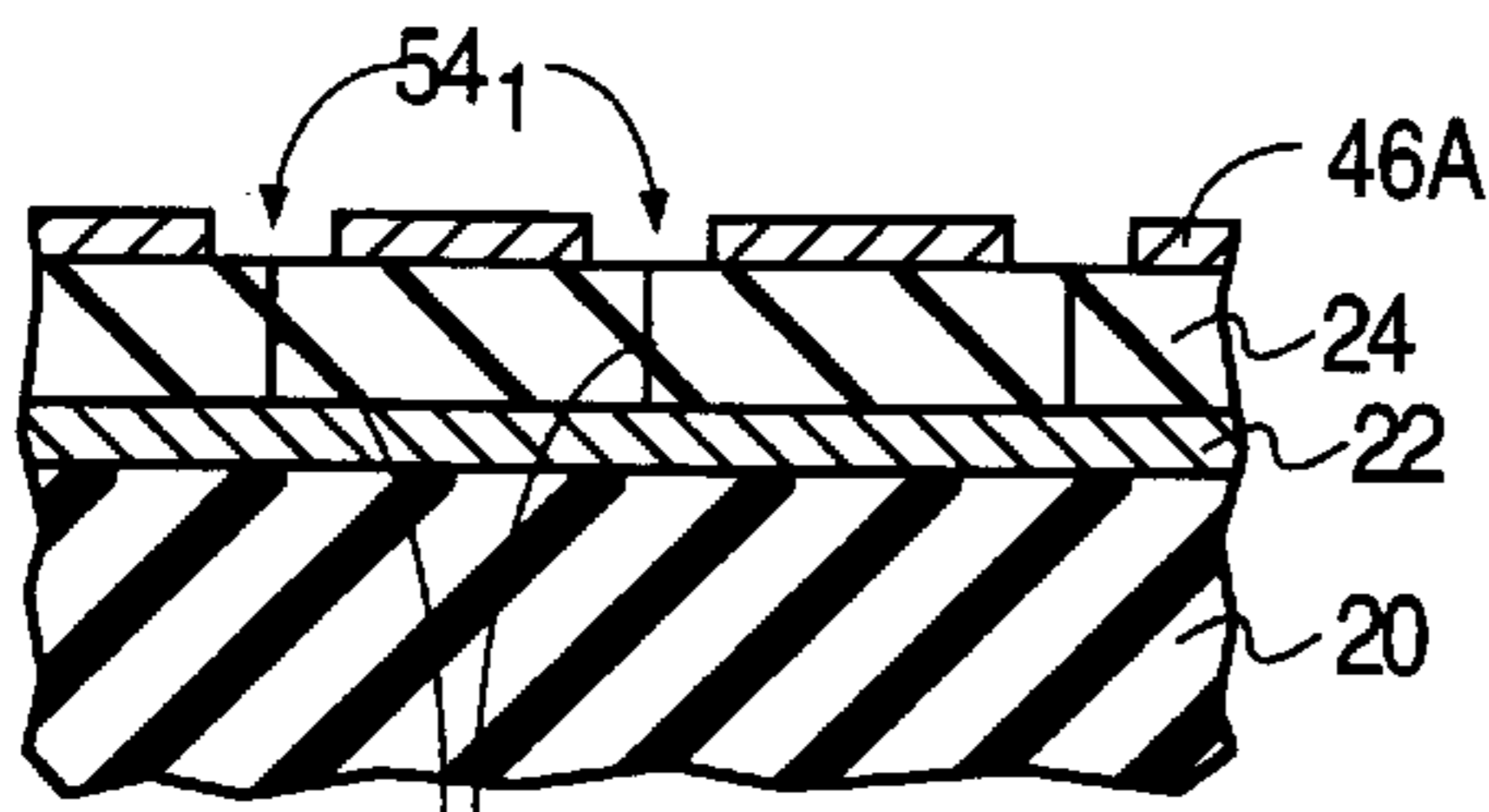


FIG. 9h



50A₁ FIG. 10a

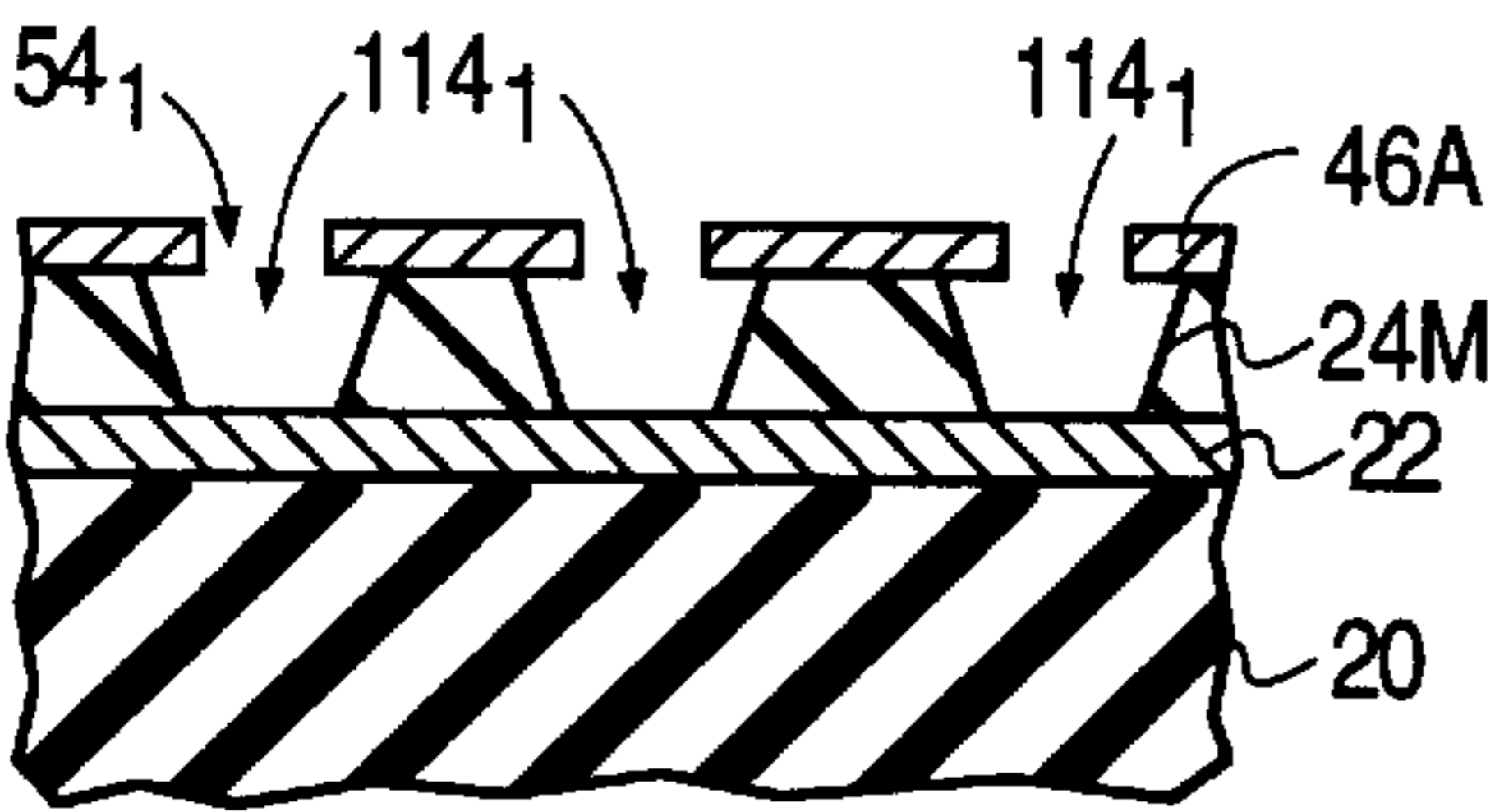


FIG. 10b

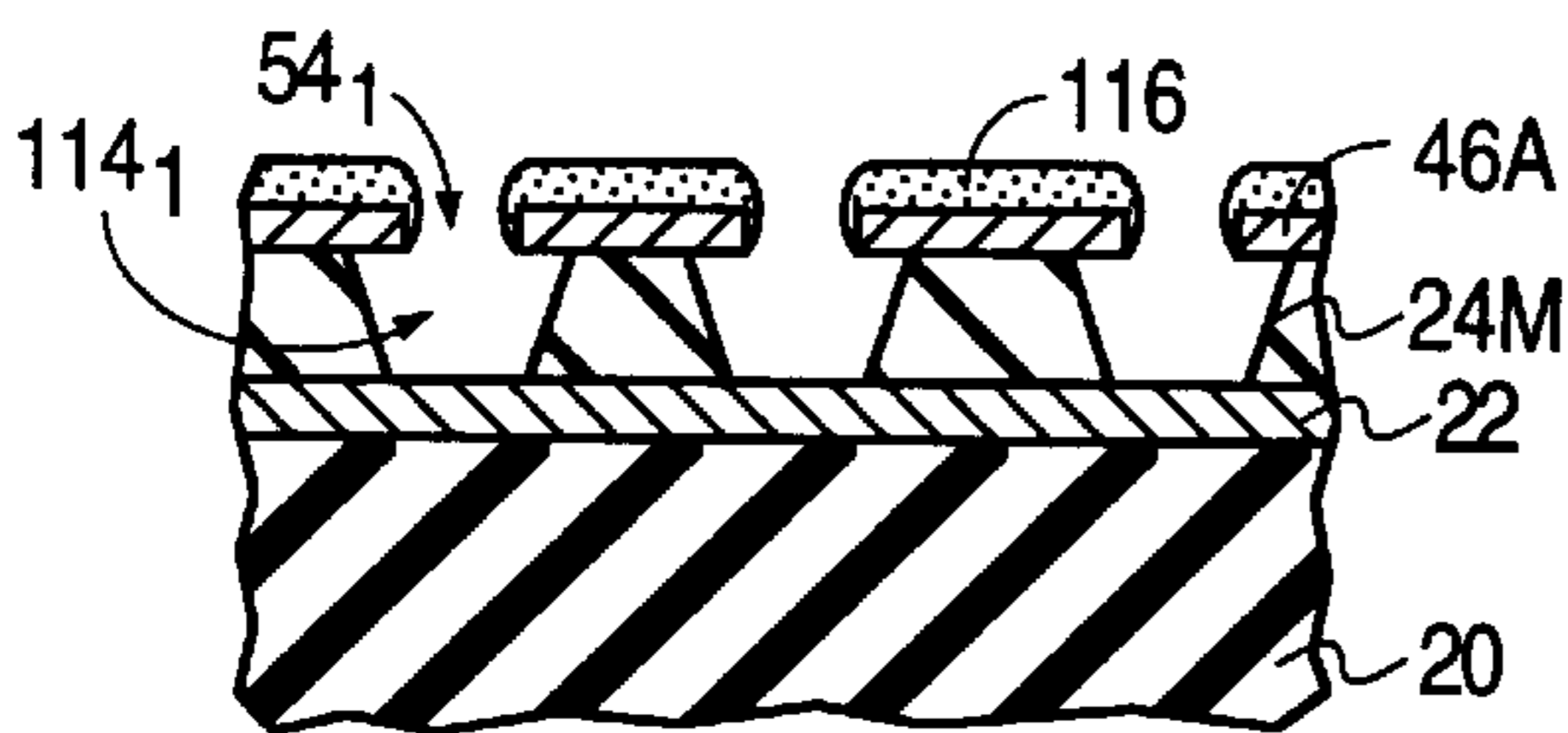


FIG. 10c

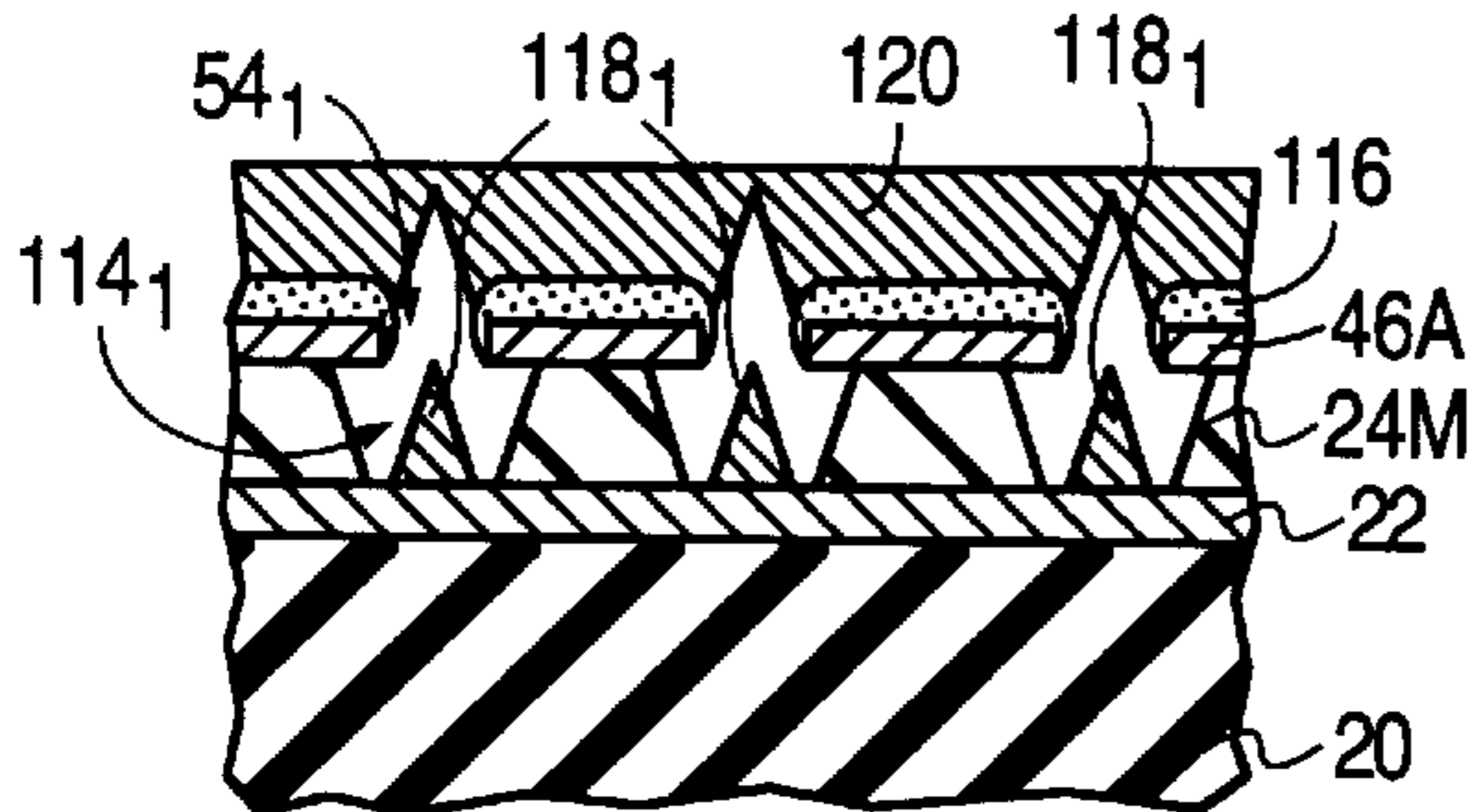


FIG. 10d

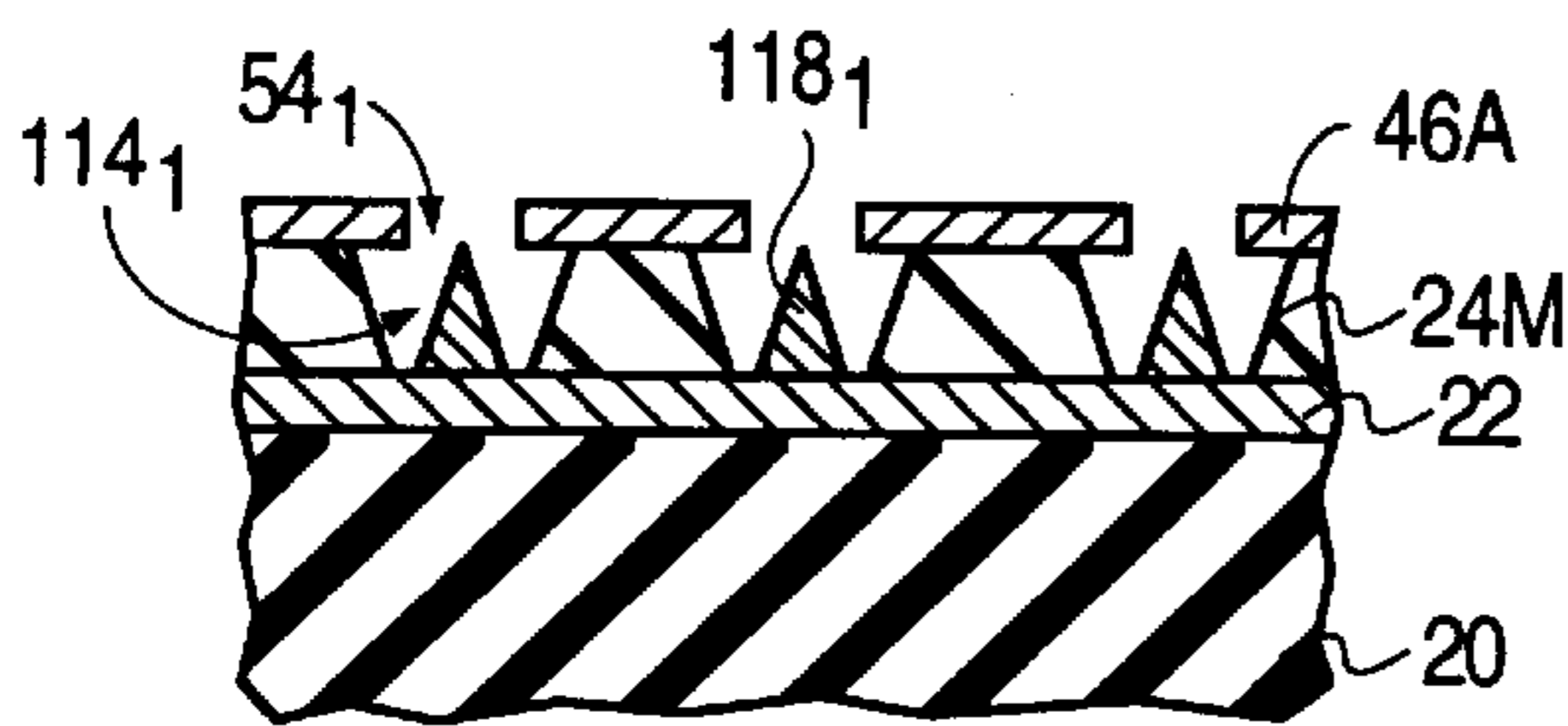


FIG. 10e

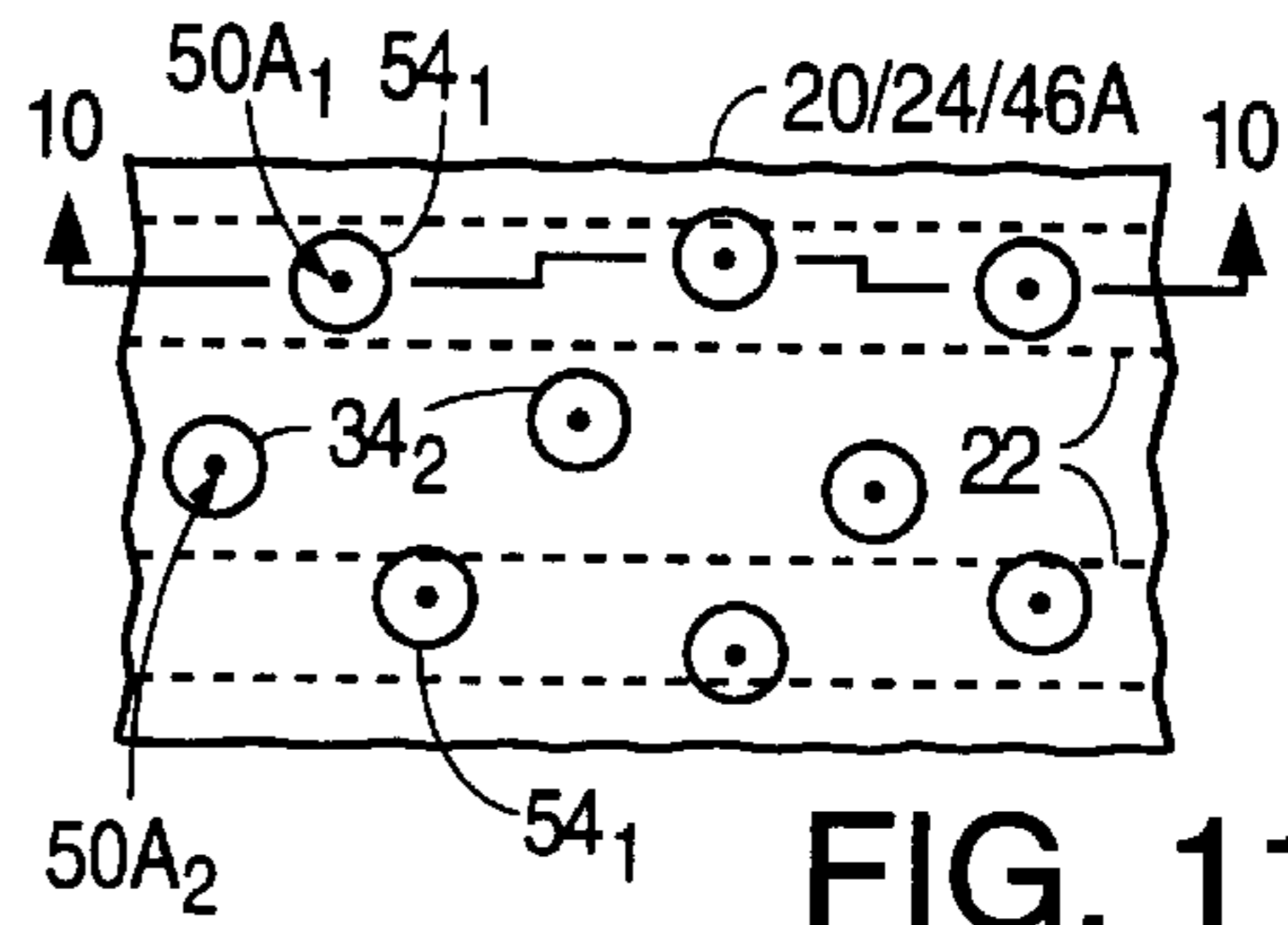


FIG. 11a

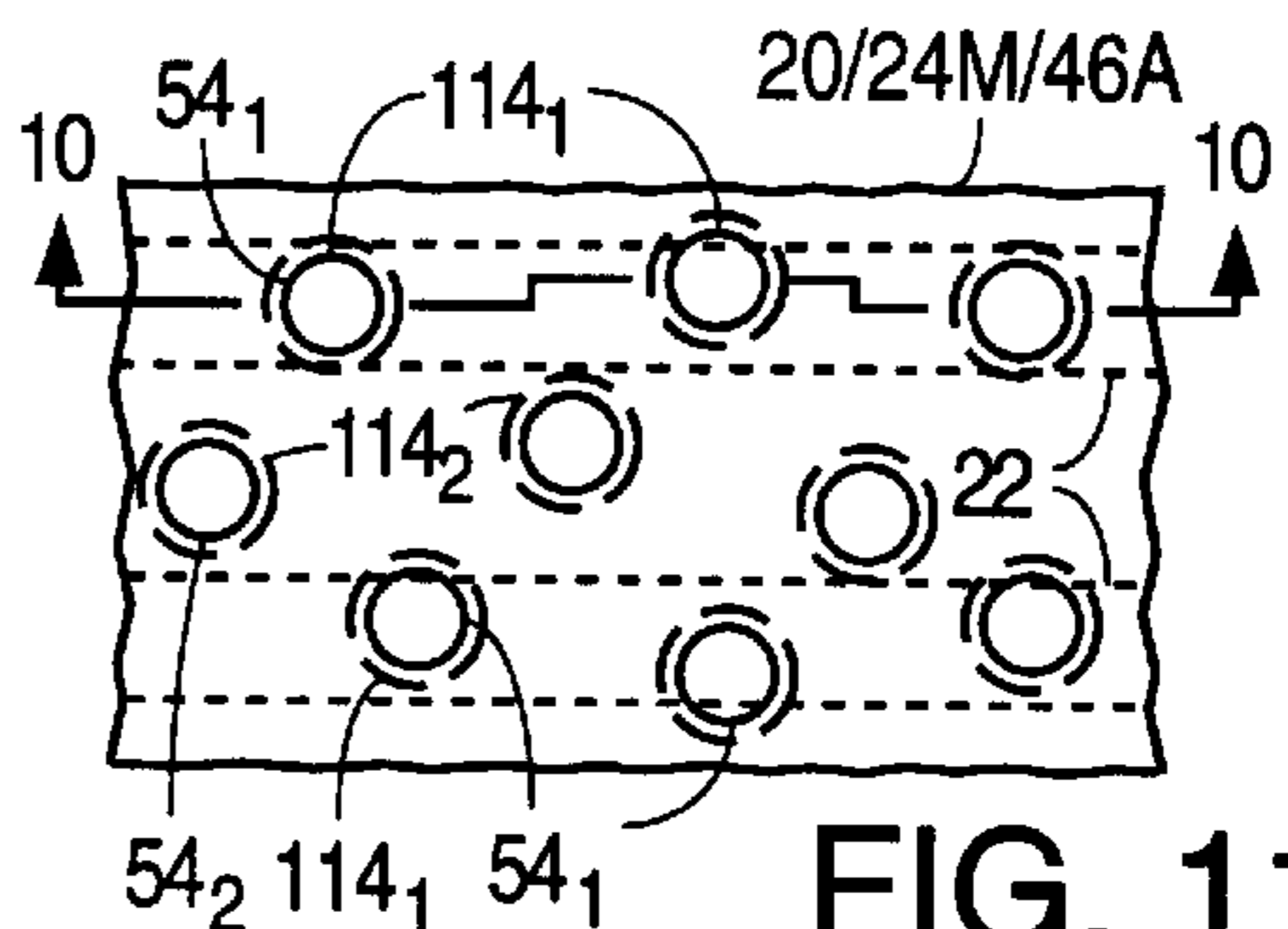


FIG. 11b

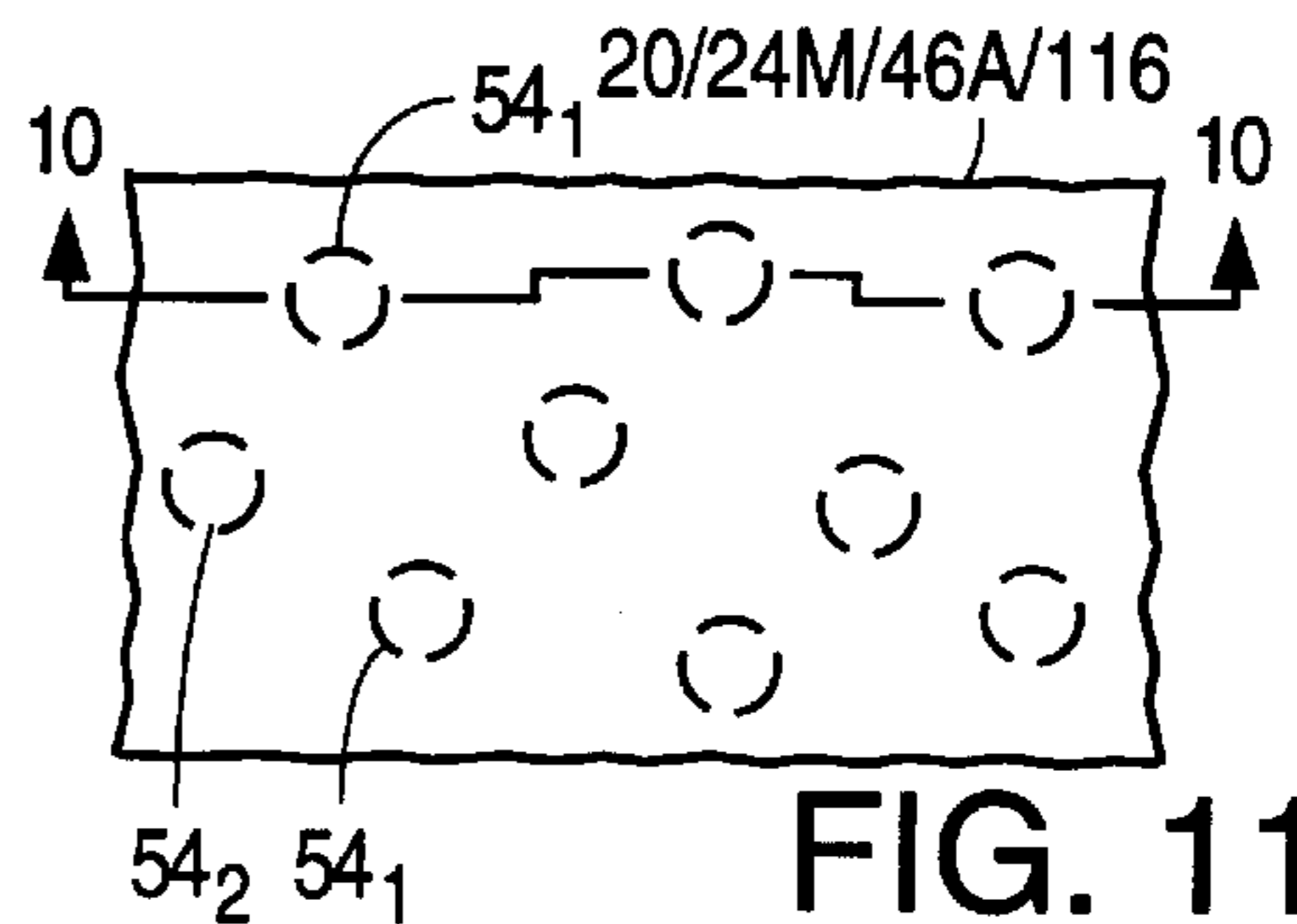


FIG. 11c

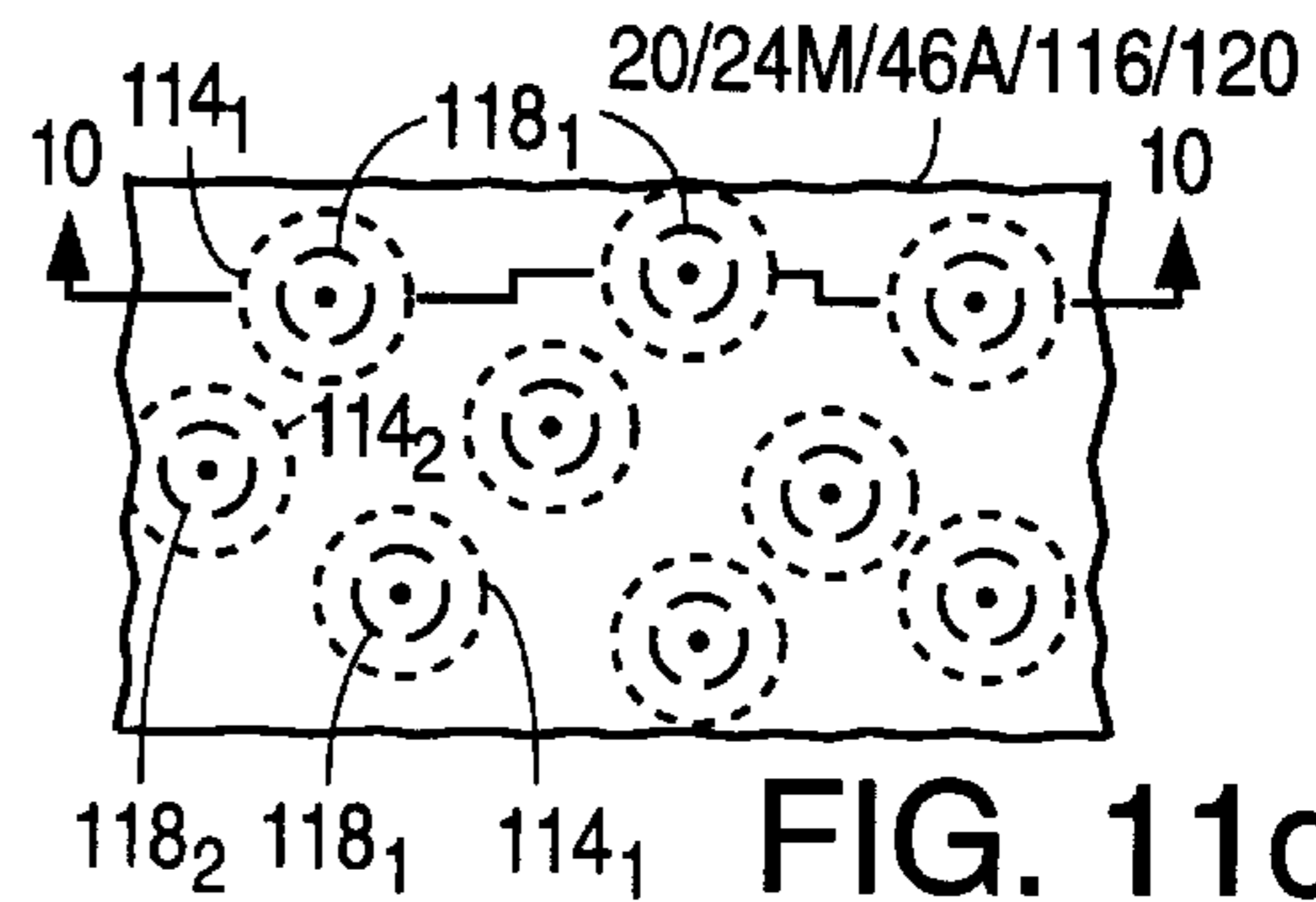


FIG. 11d

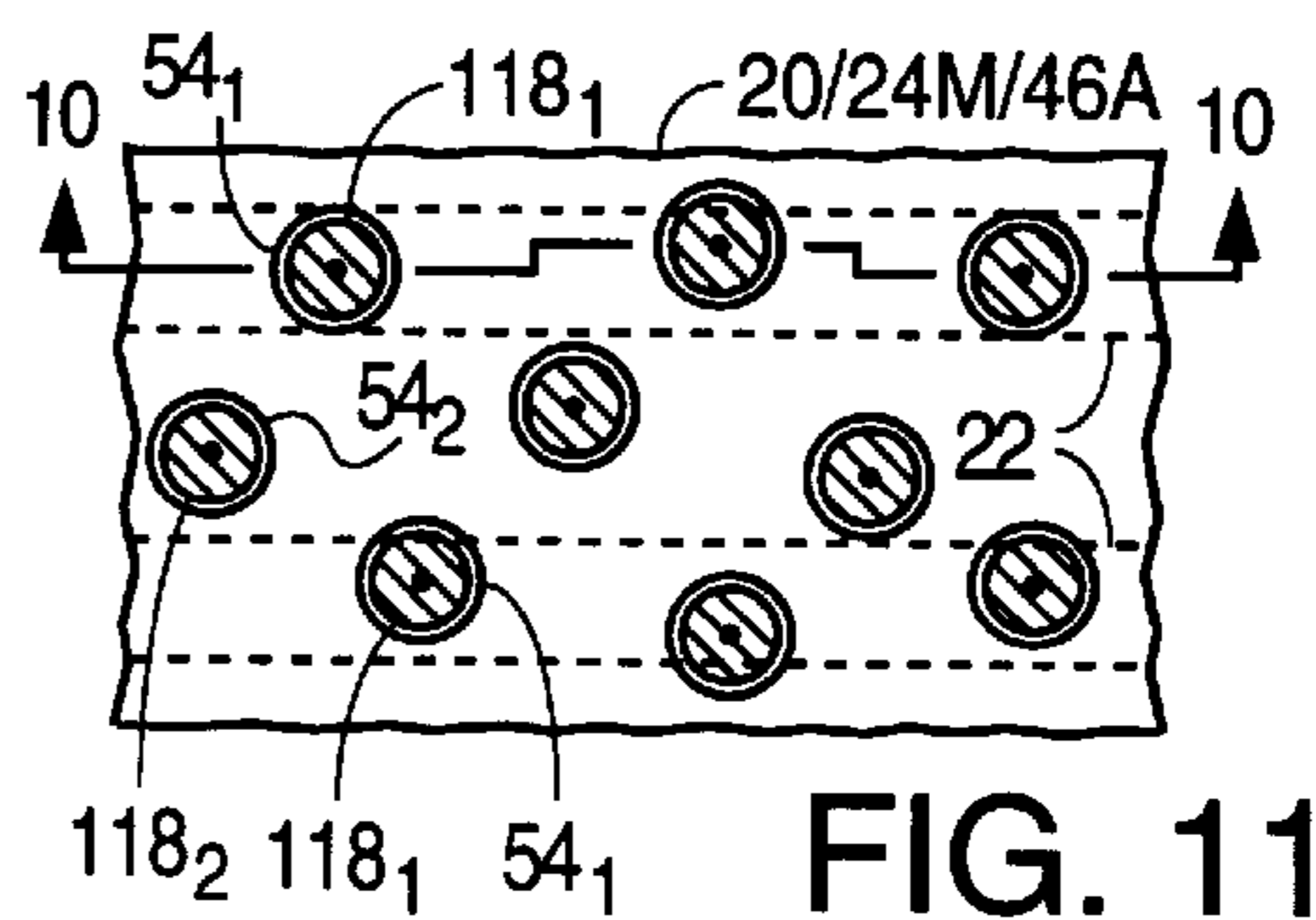


FIG. 11e

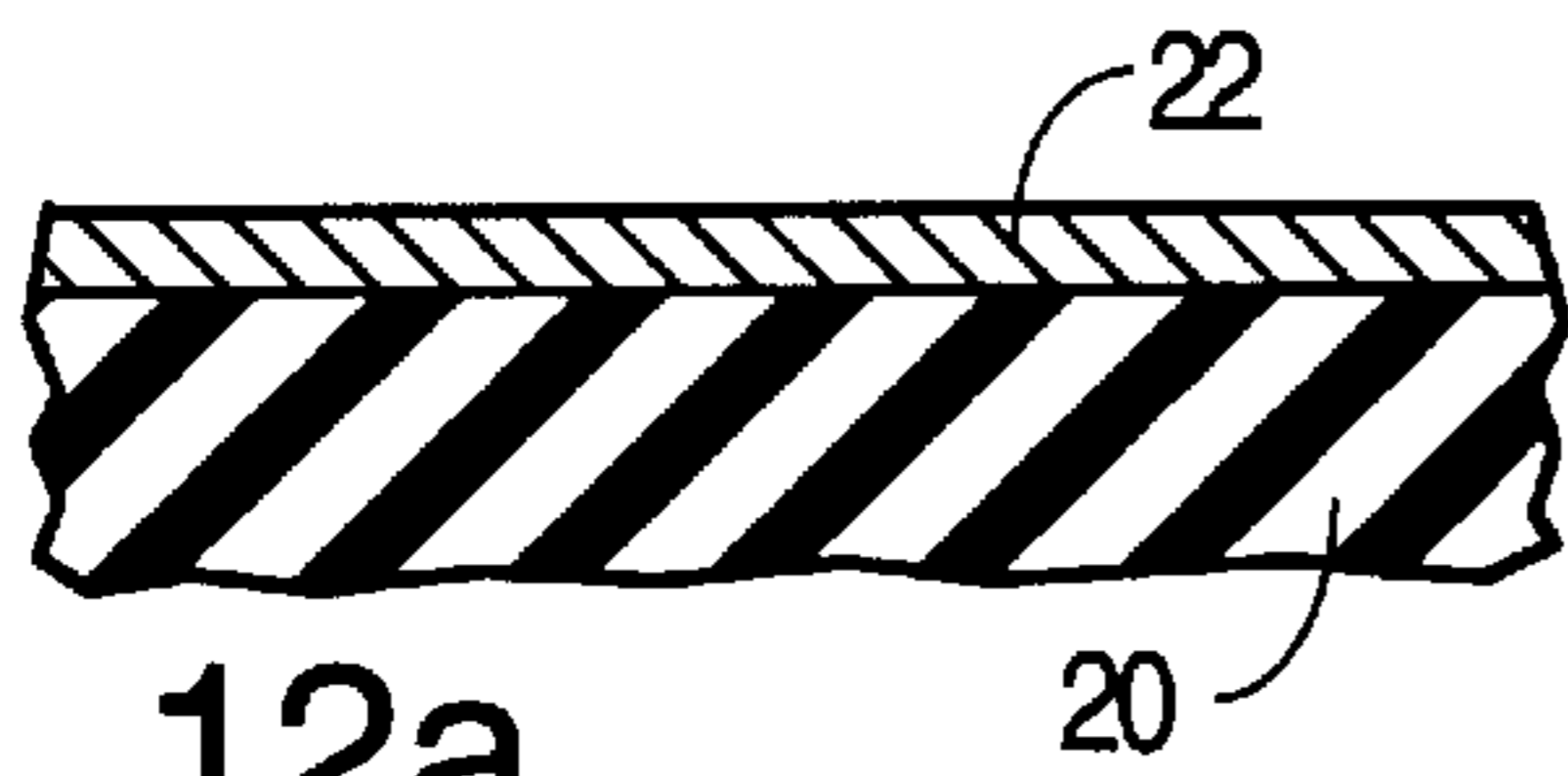


FIG. 12a

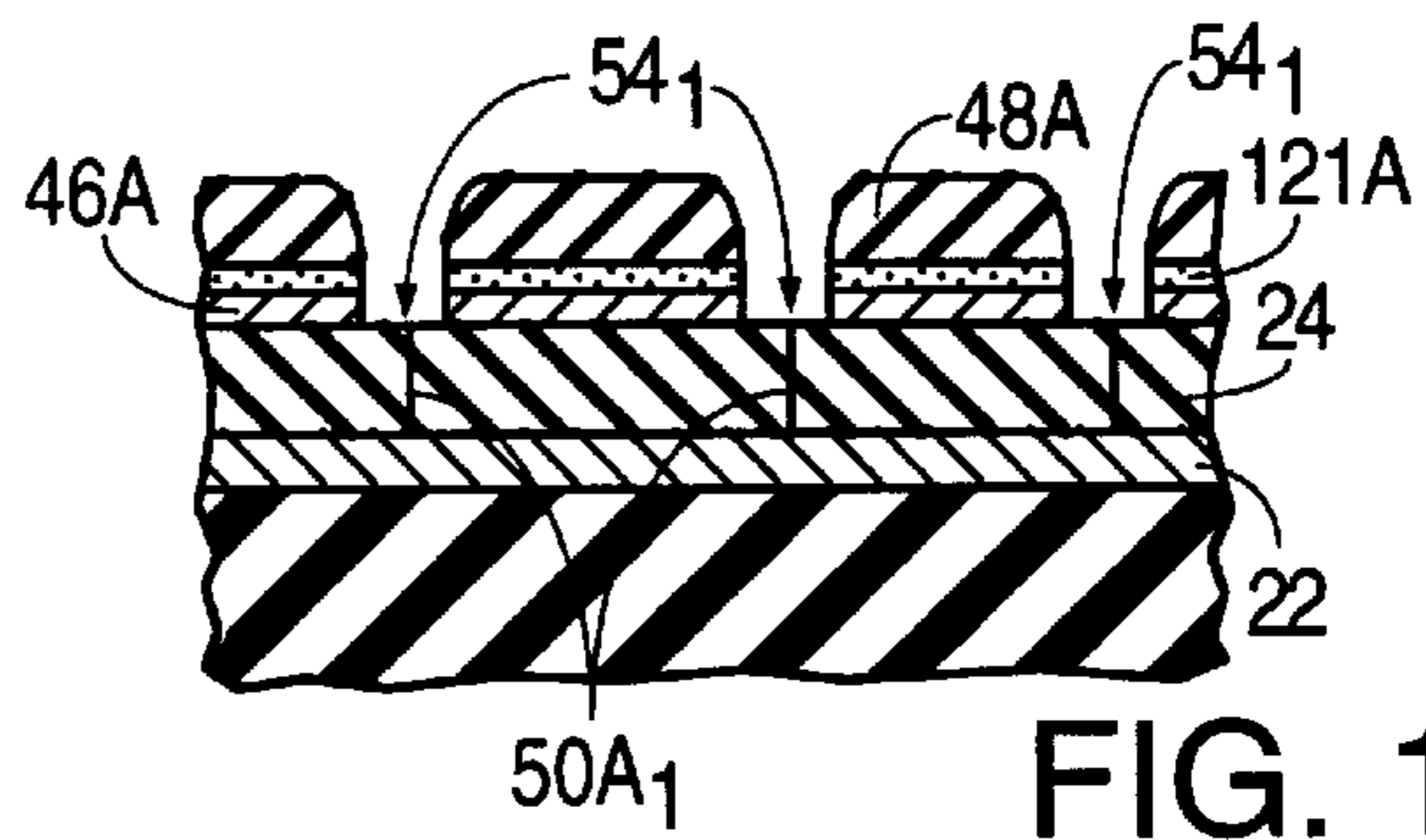


FIG. 12f

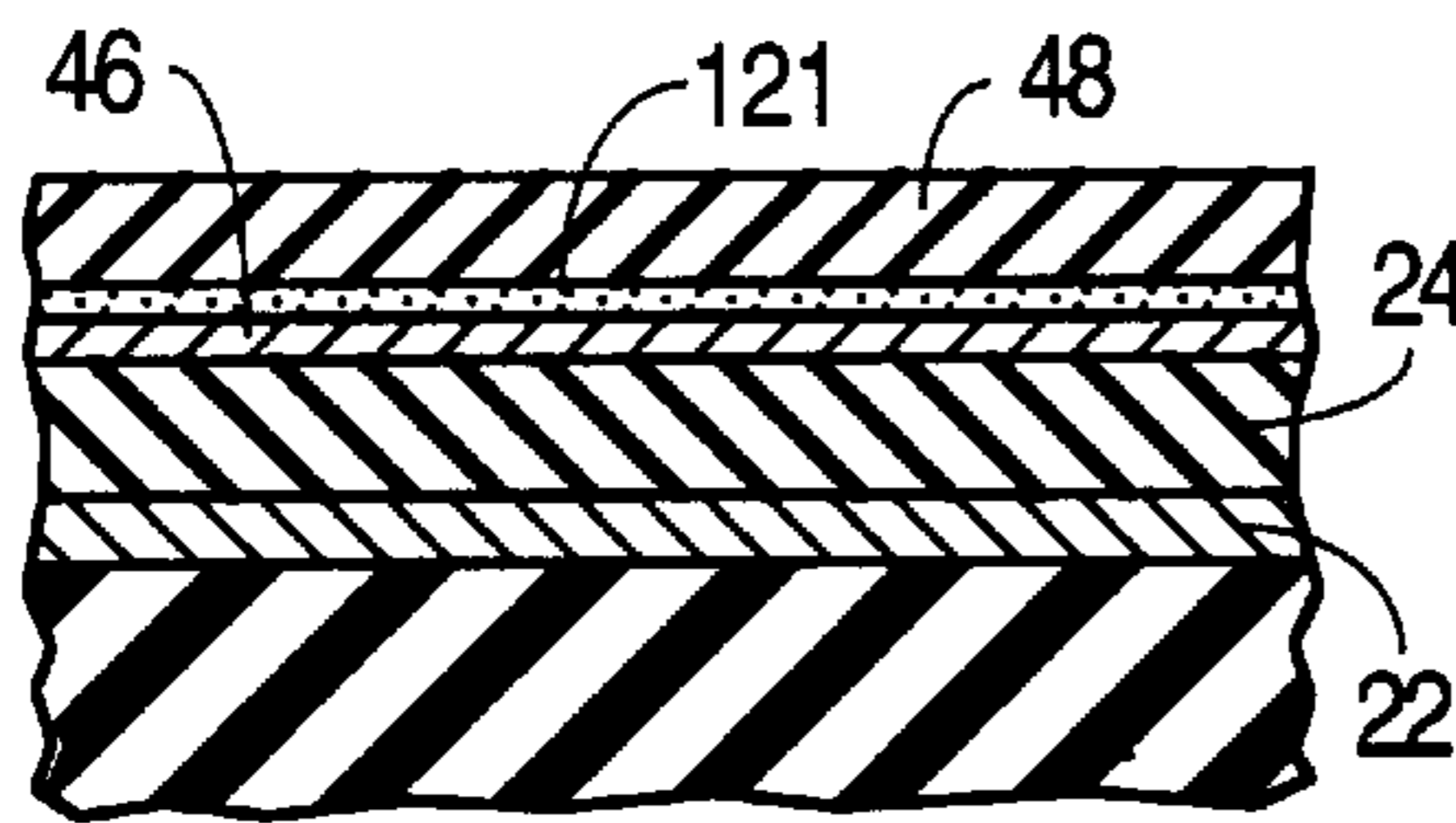


FIG. 12b

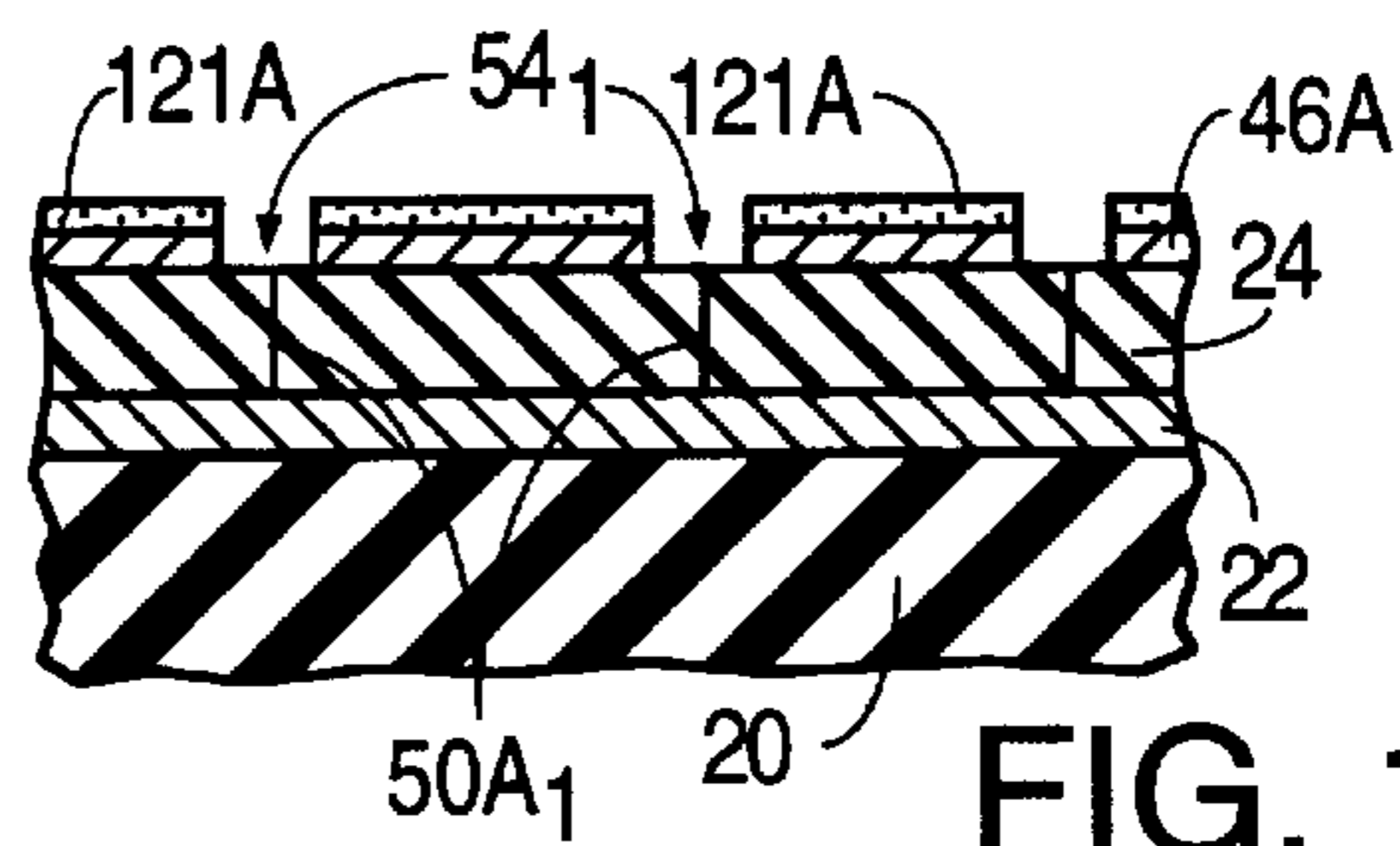


FIG. 12g

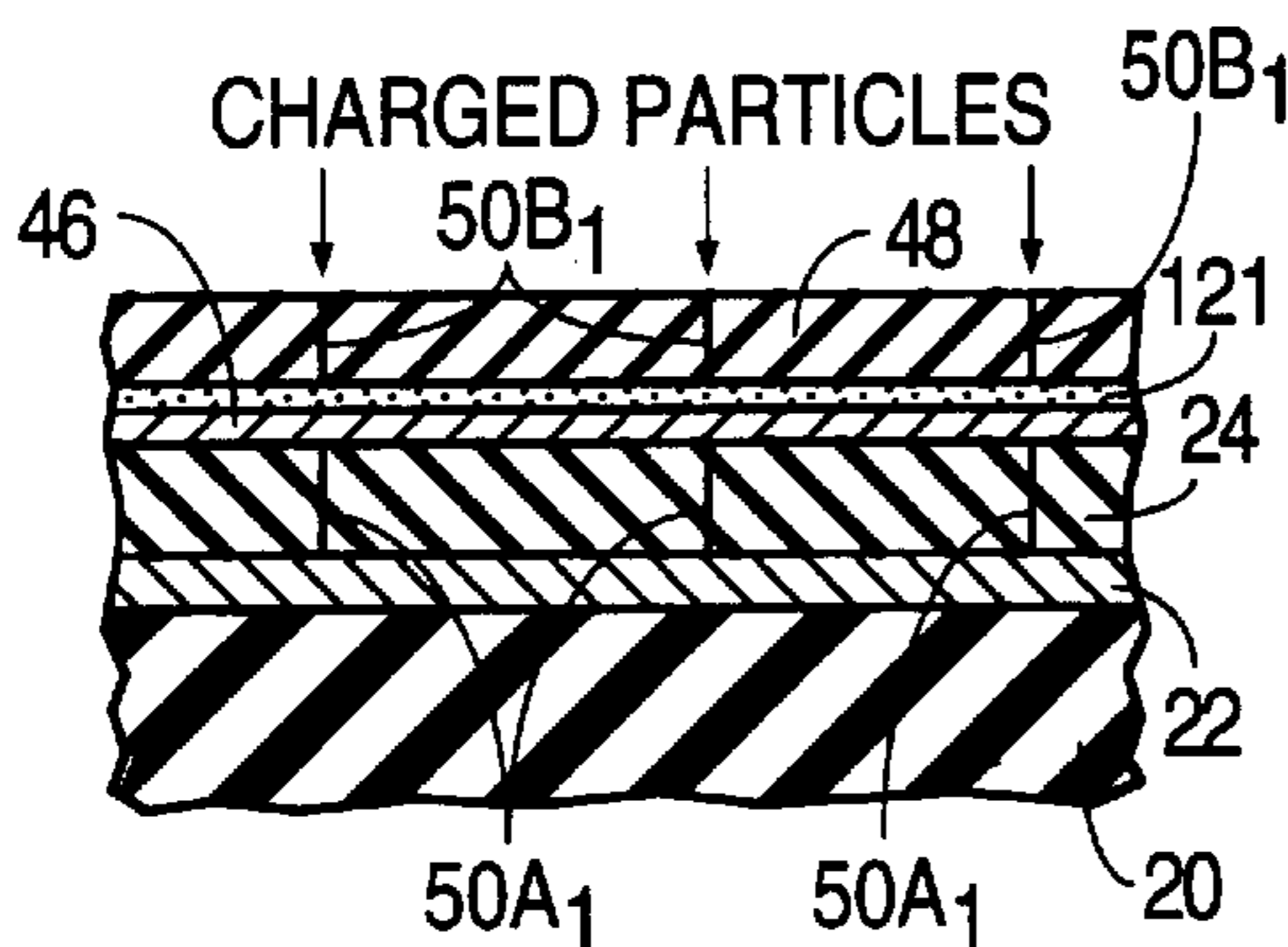


FIG. 12c

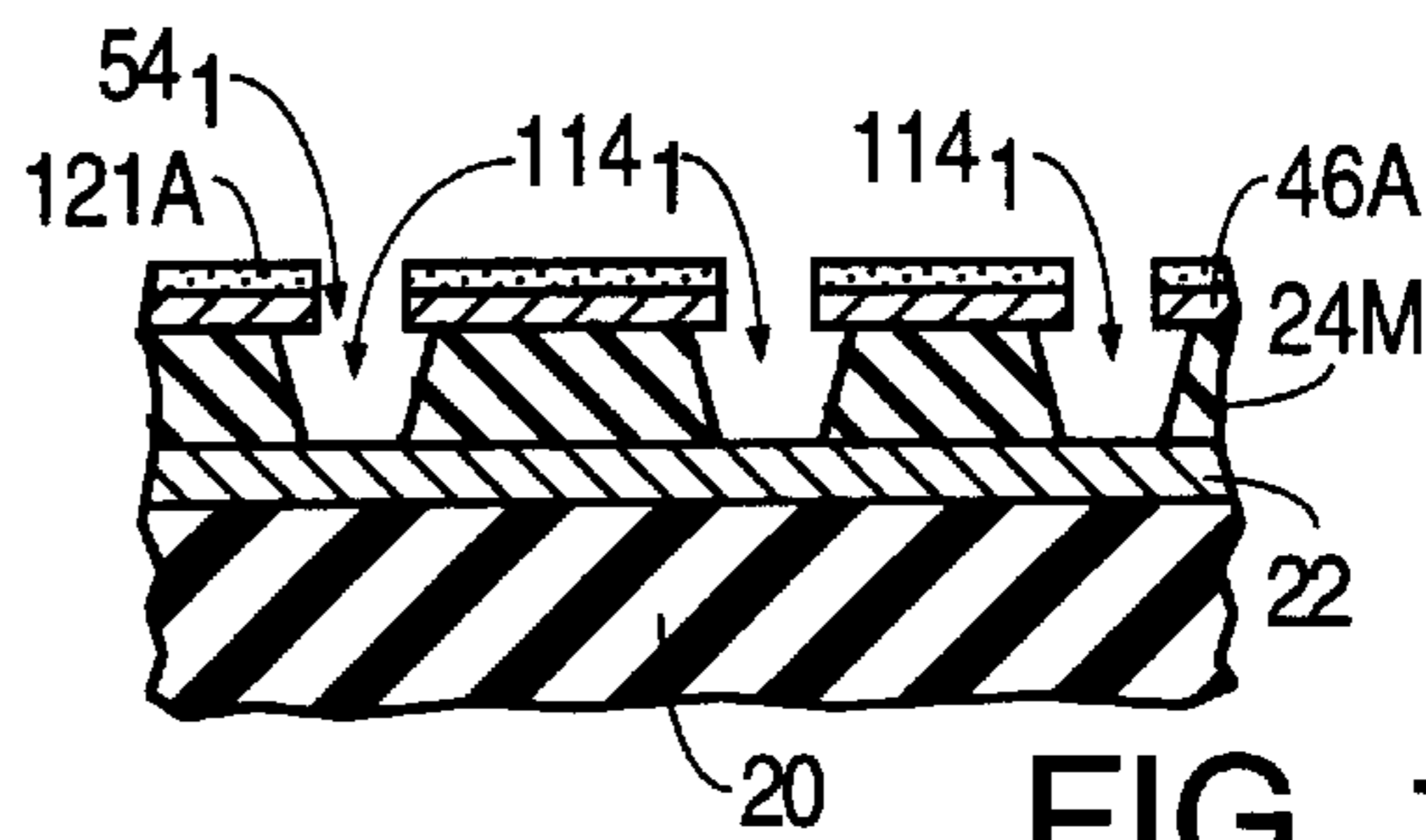


FIG. 12h

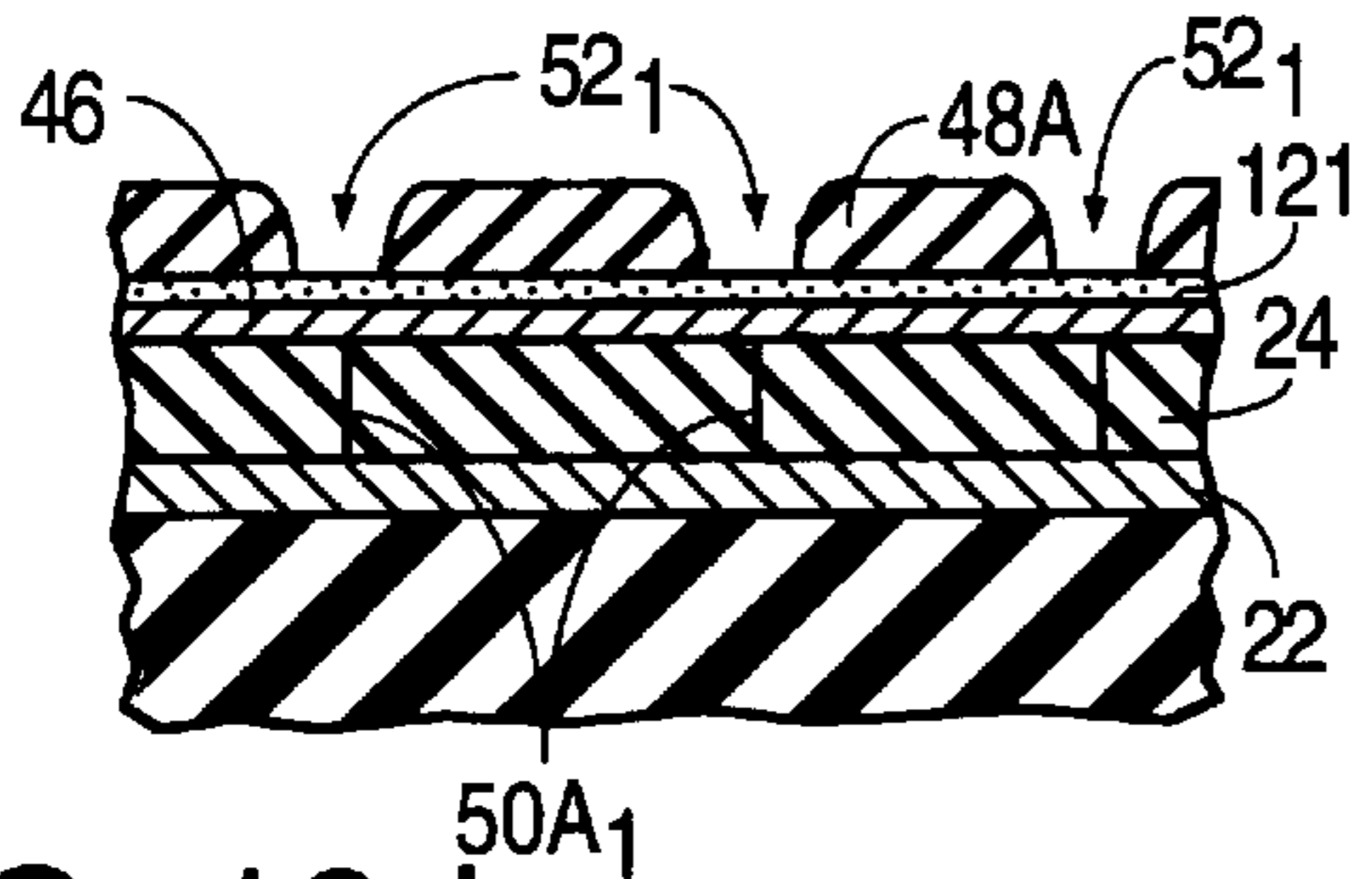


FIG. 12d

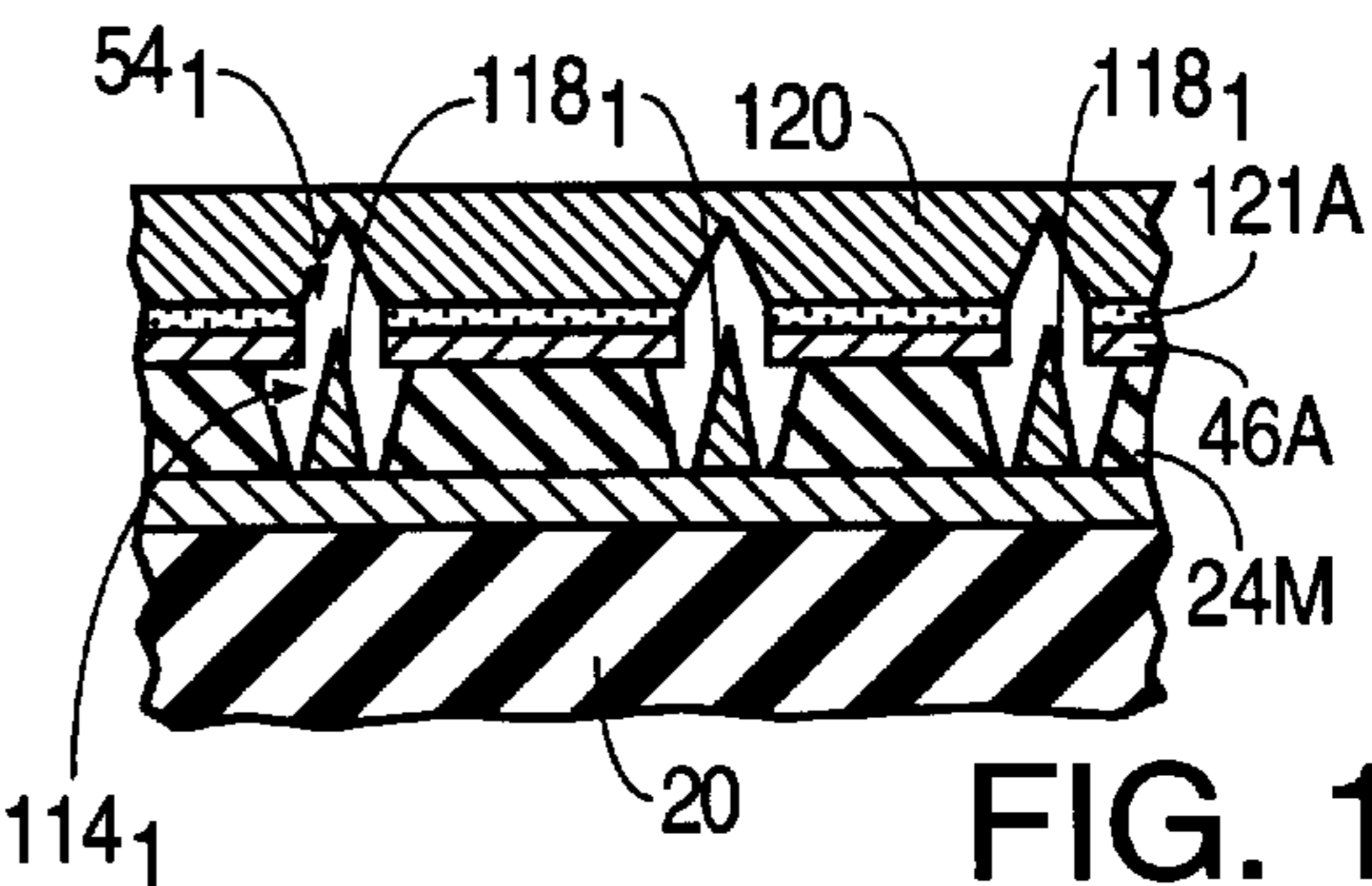


FIG. 12i

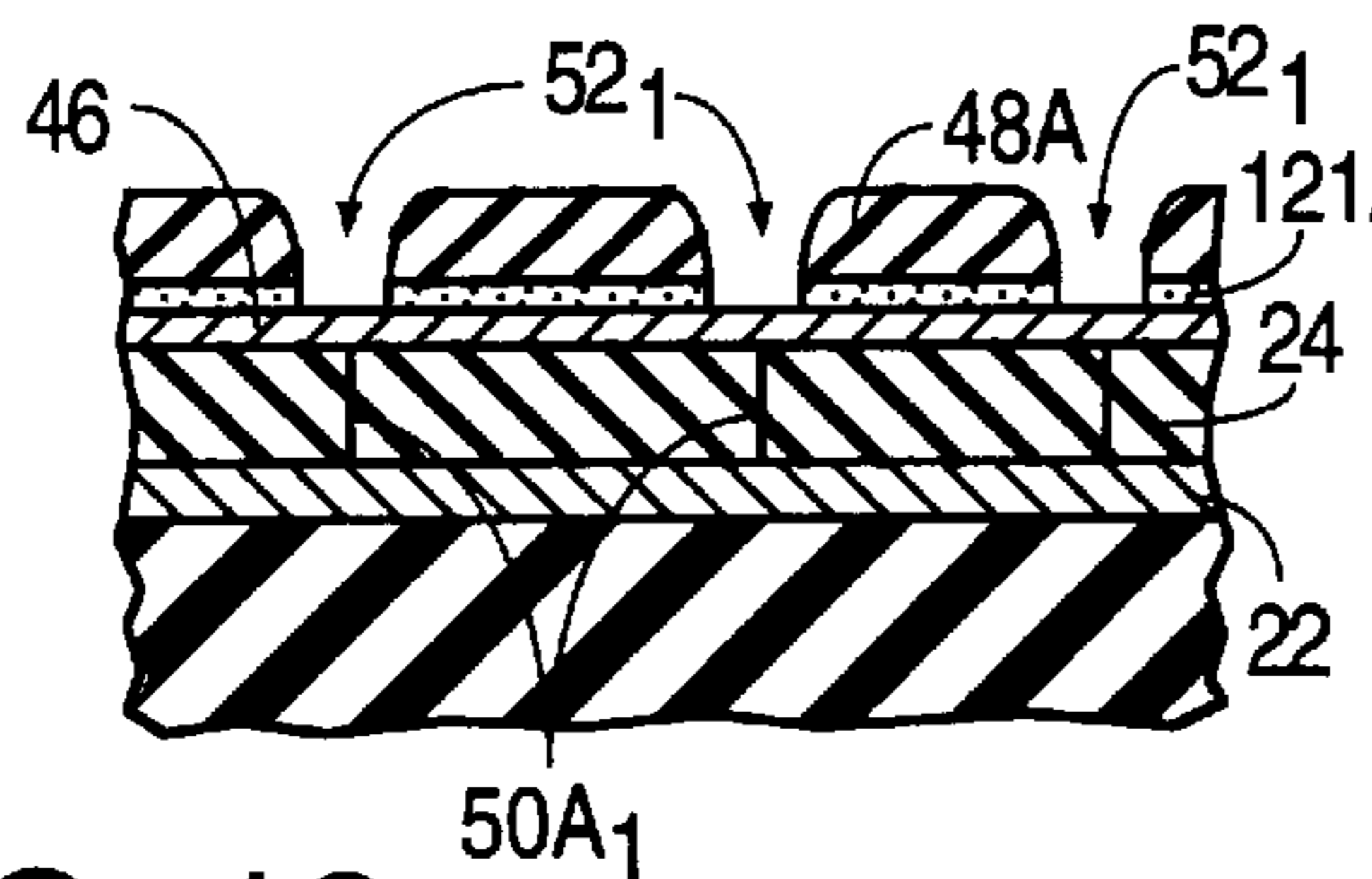


FIG. 12e

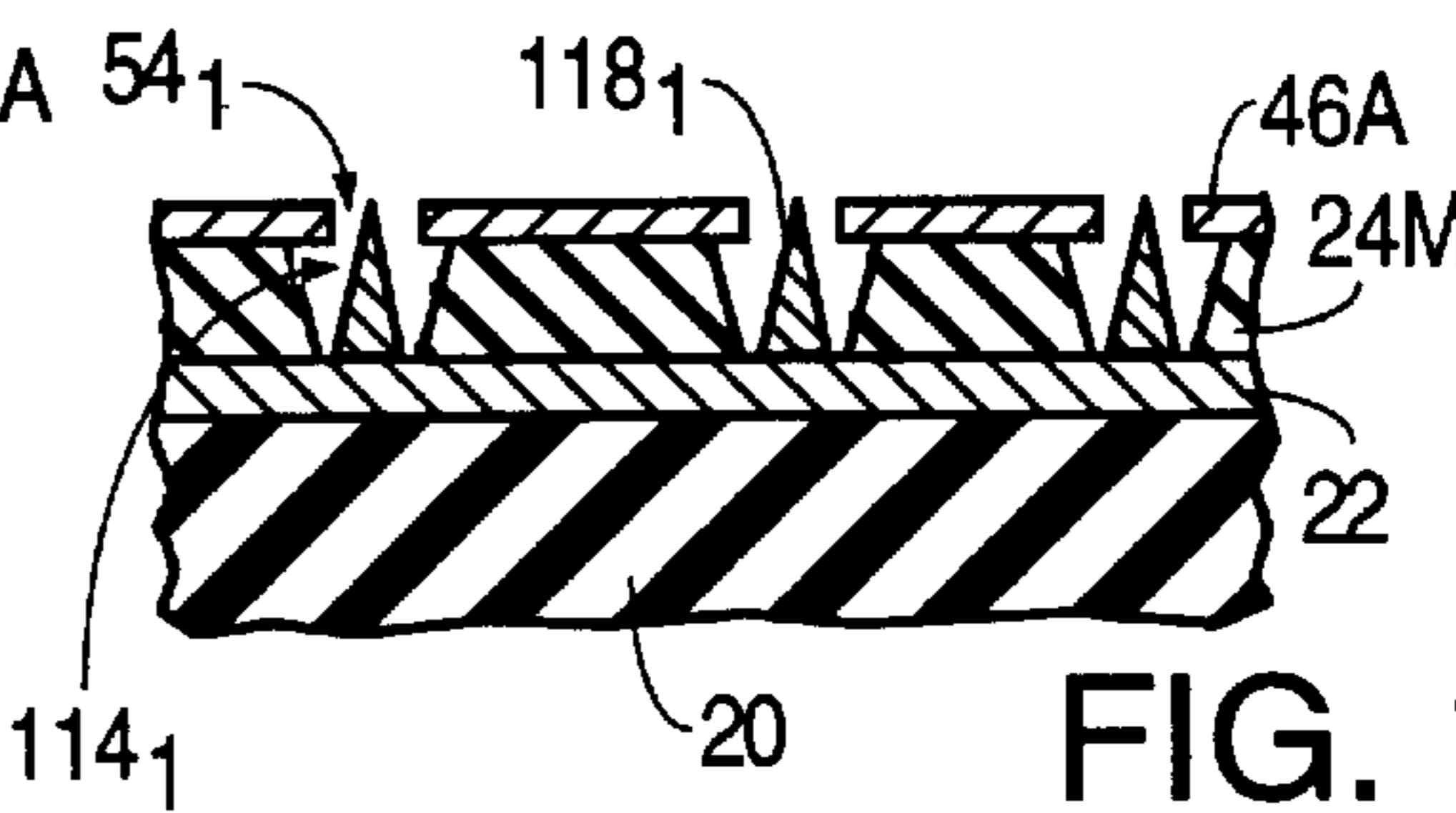


FIG. 12j

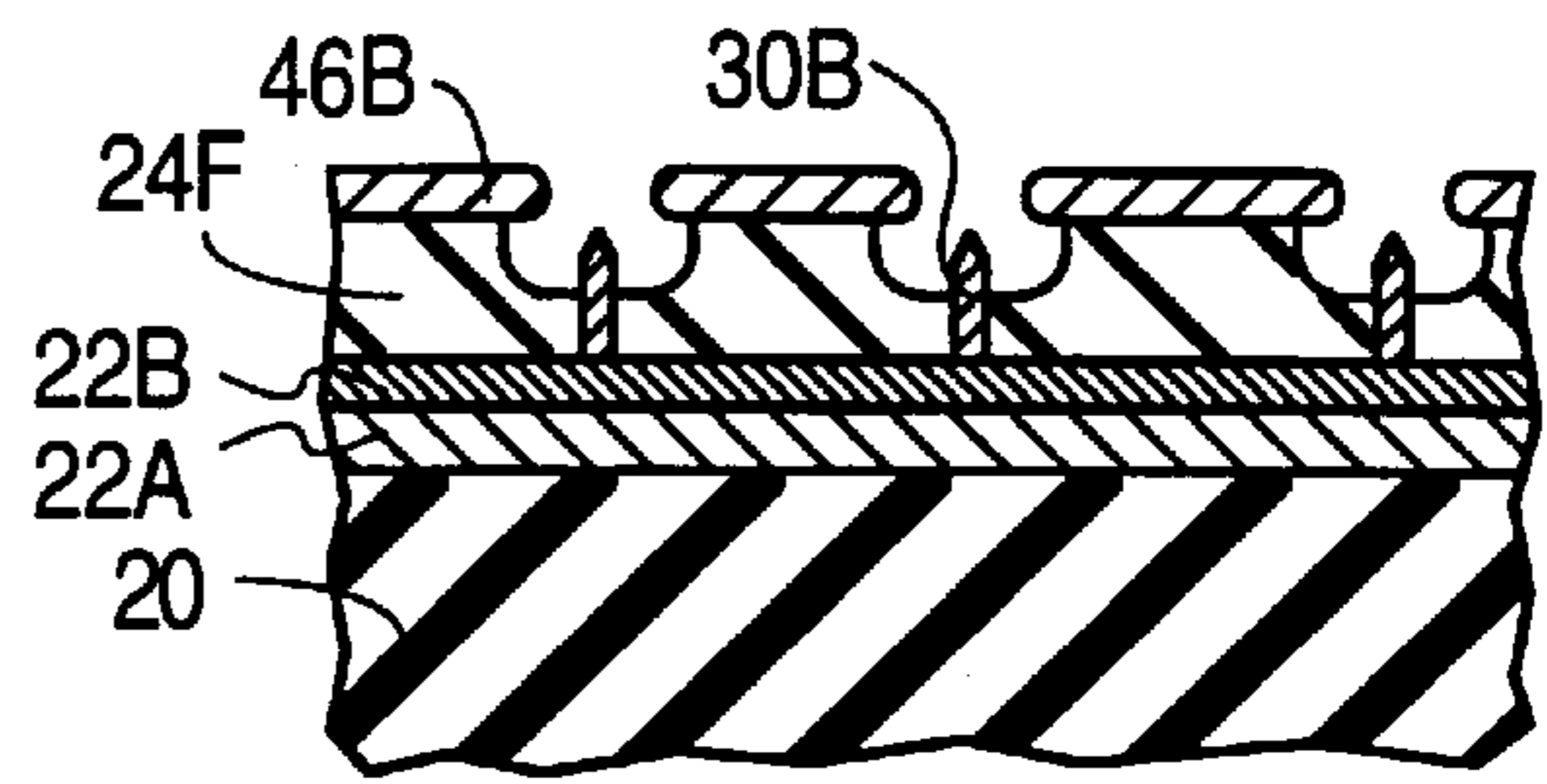


FIG. 14.1

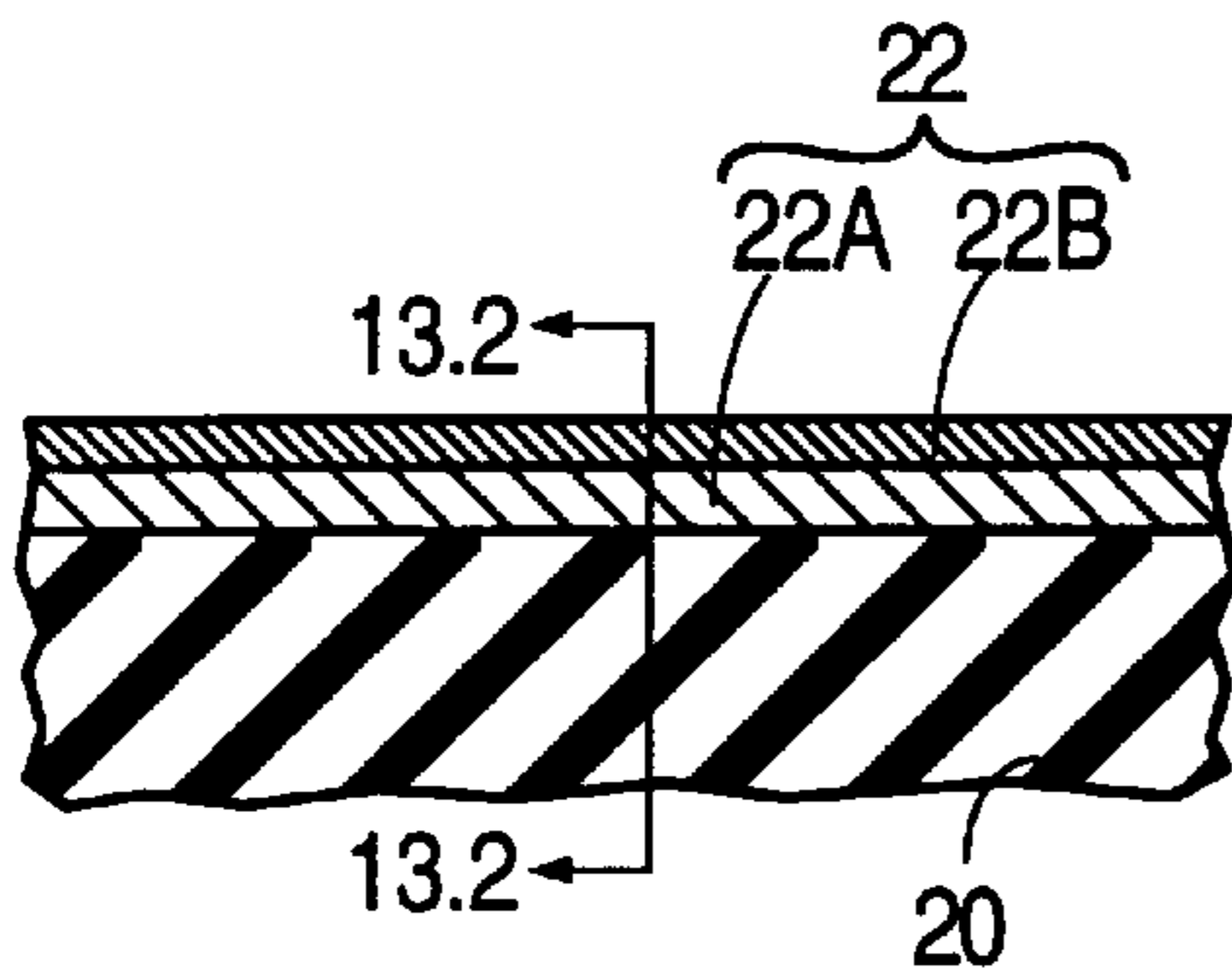


FIG. 13.1

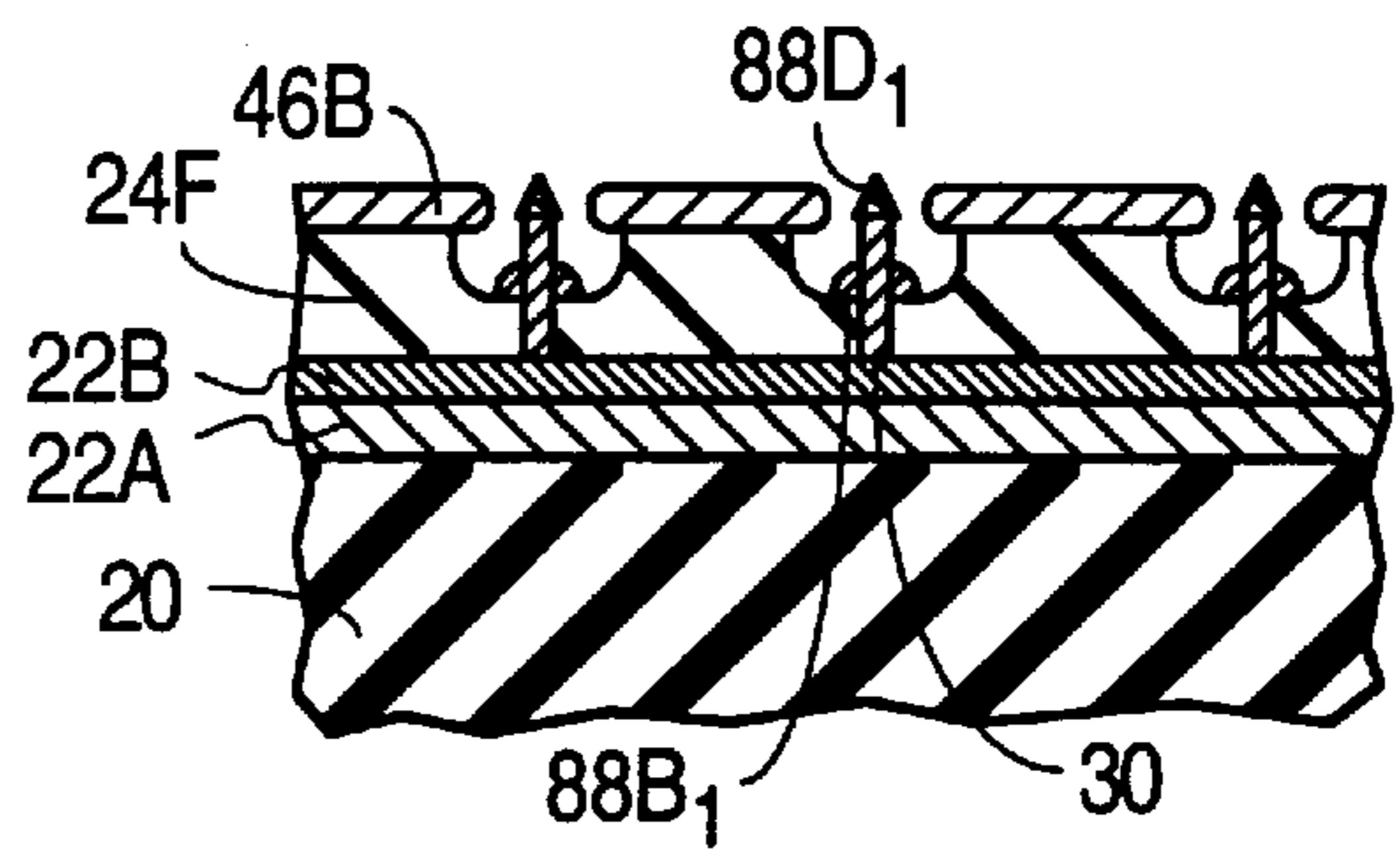


FIG. 14.2

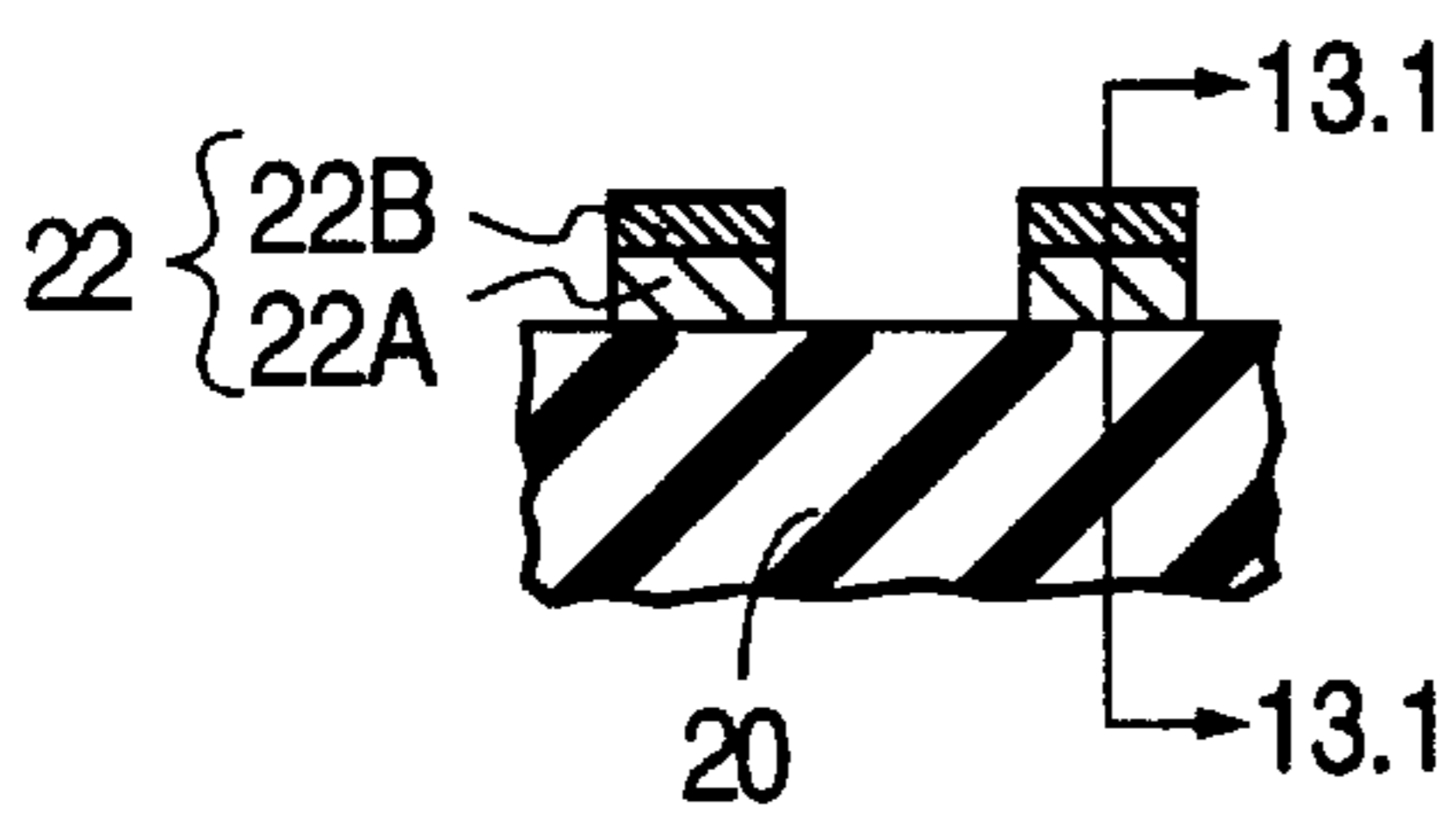


FIG. 13.2

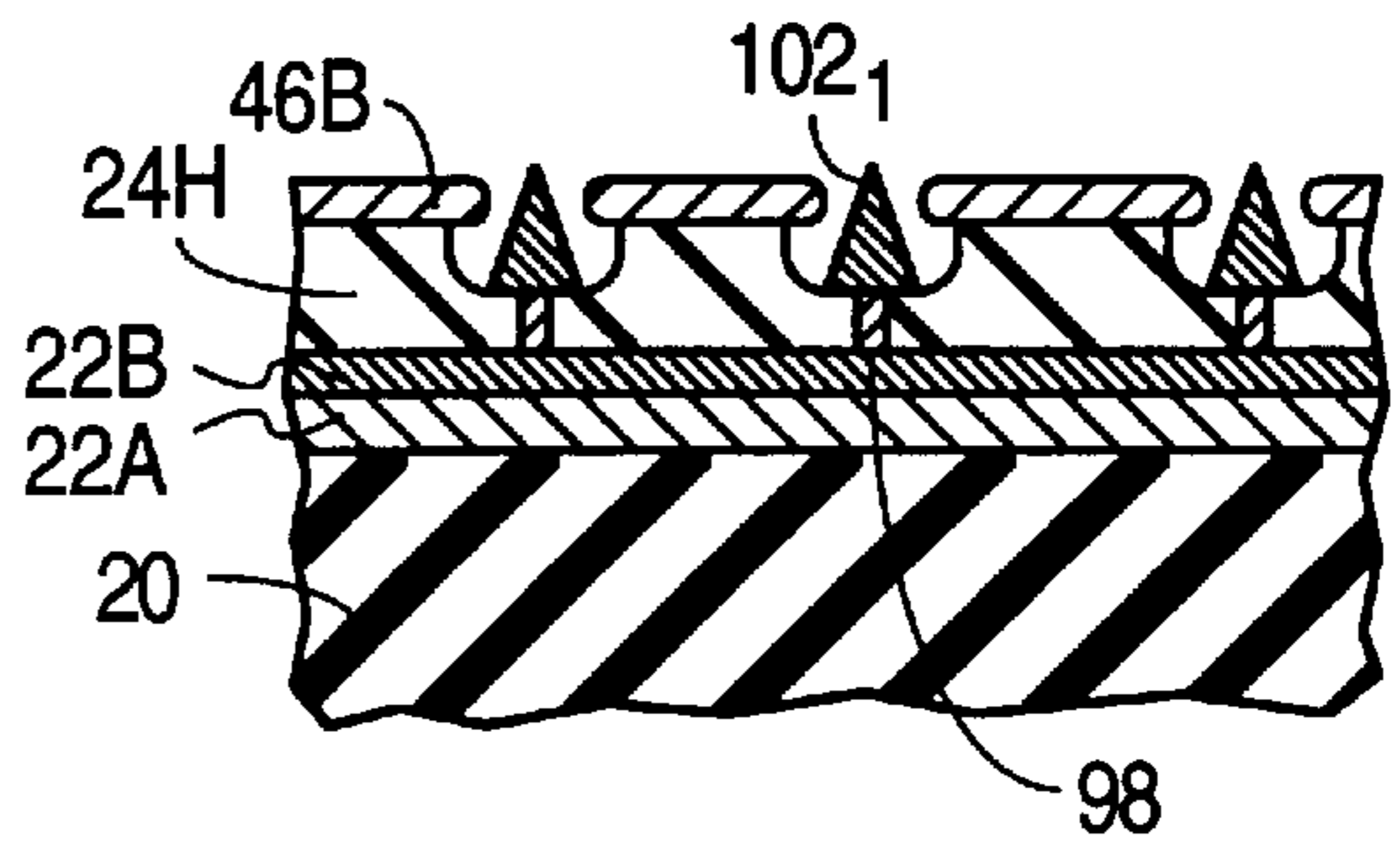


FIG. 14.3

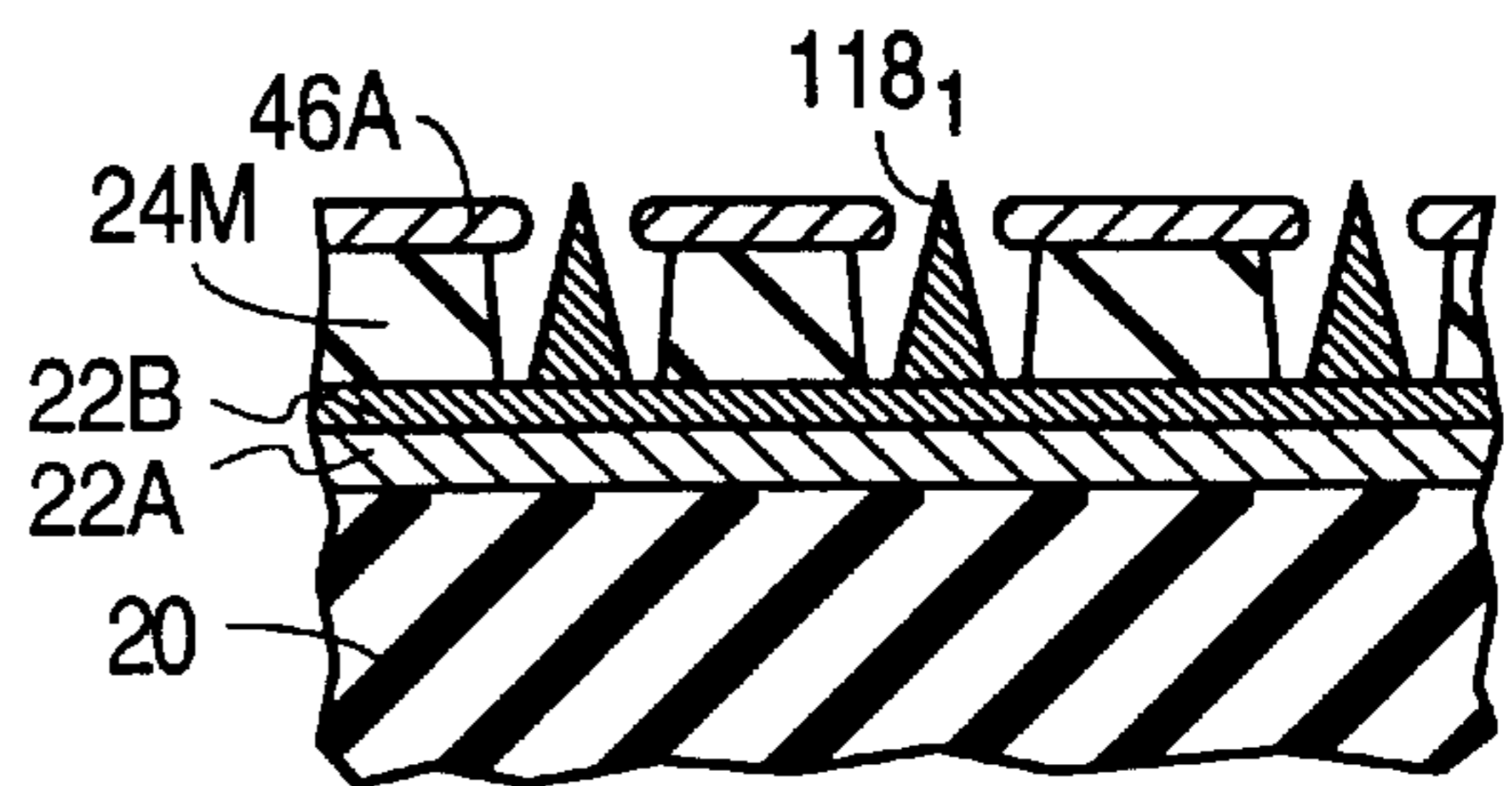


FIG. 14.4

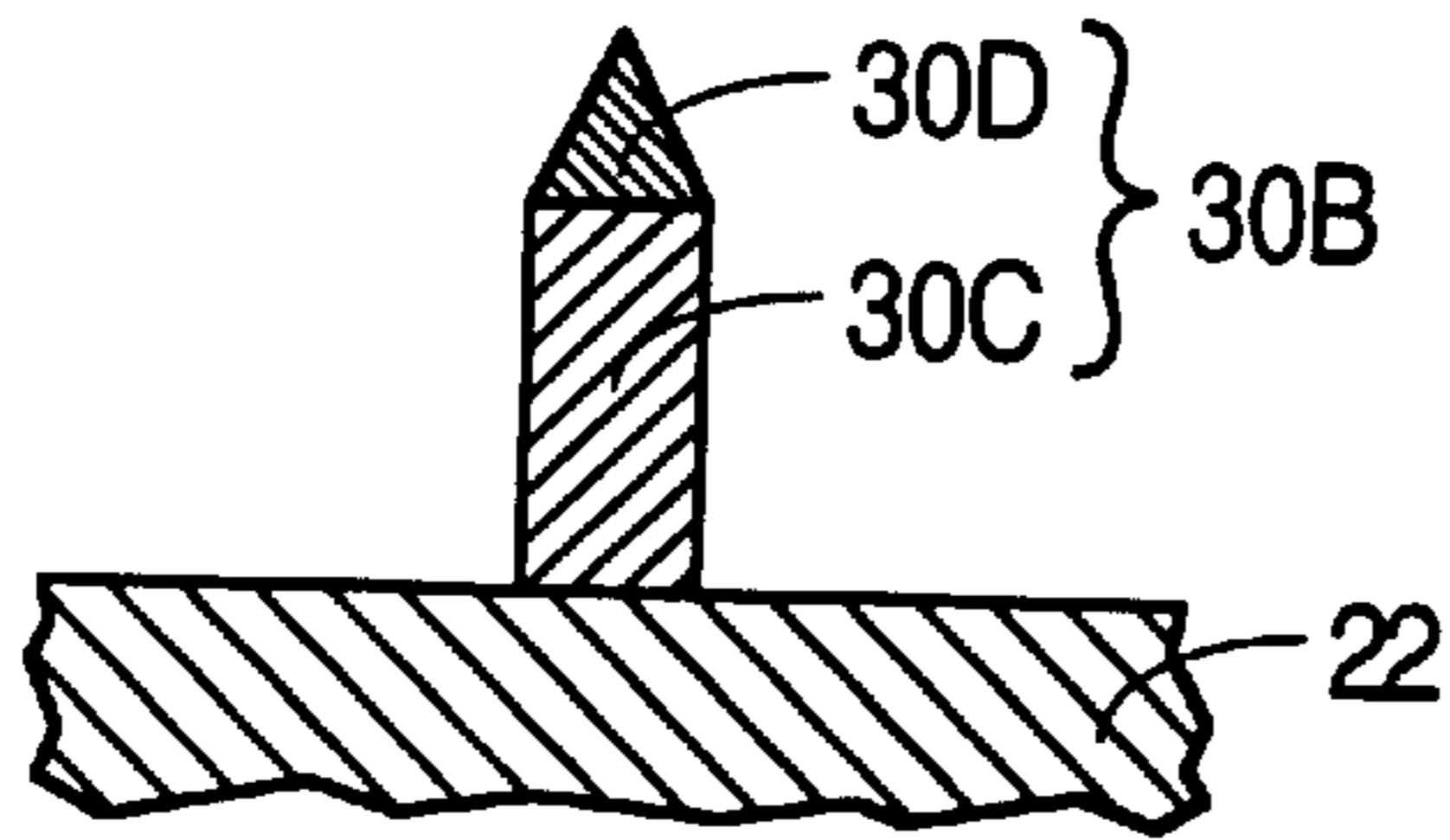


FIG. 15.1

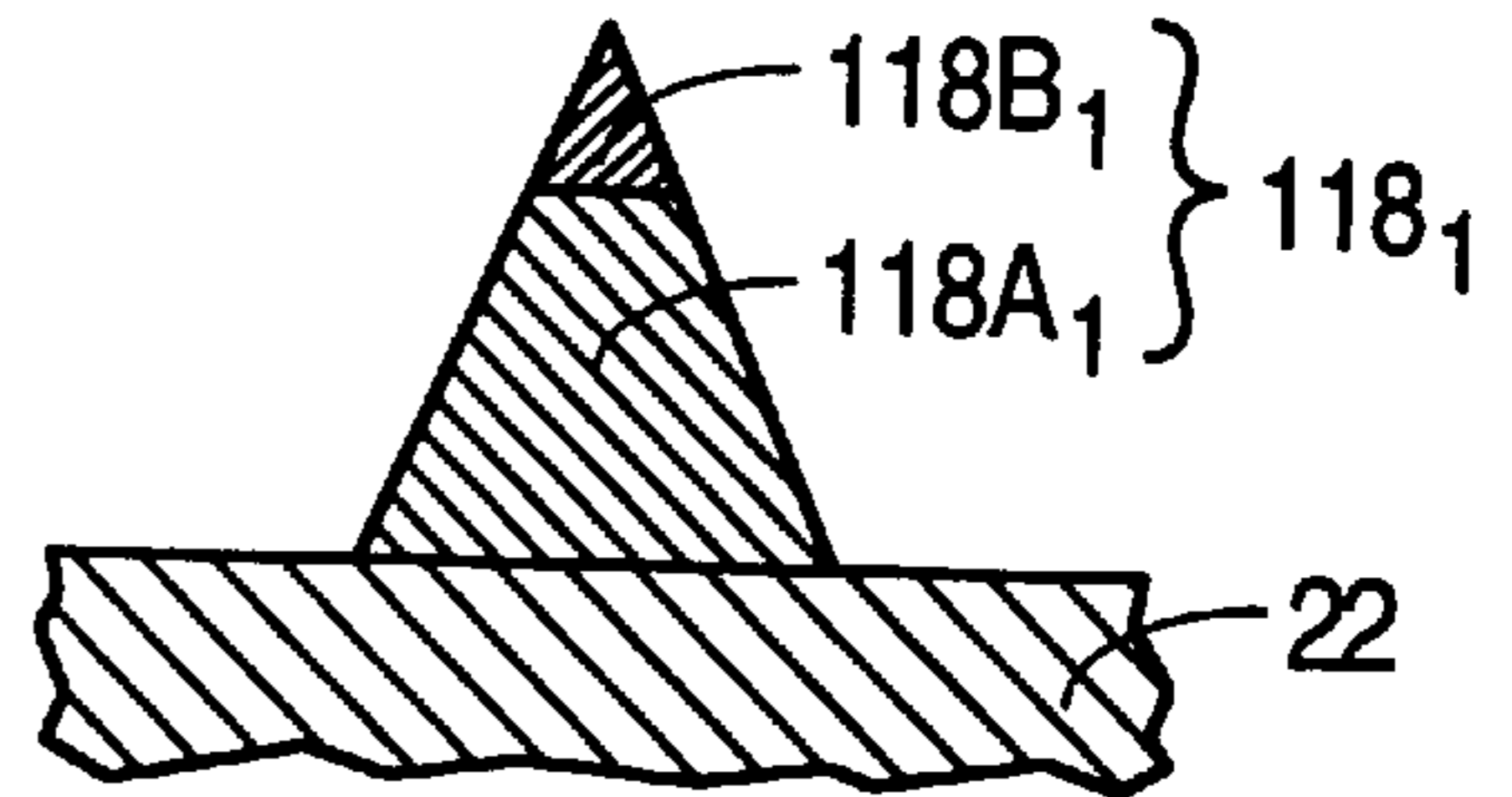


FIG. 15.2

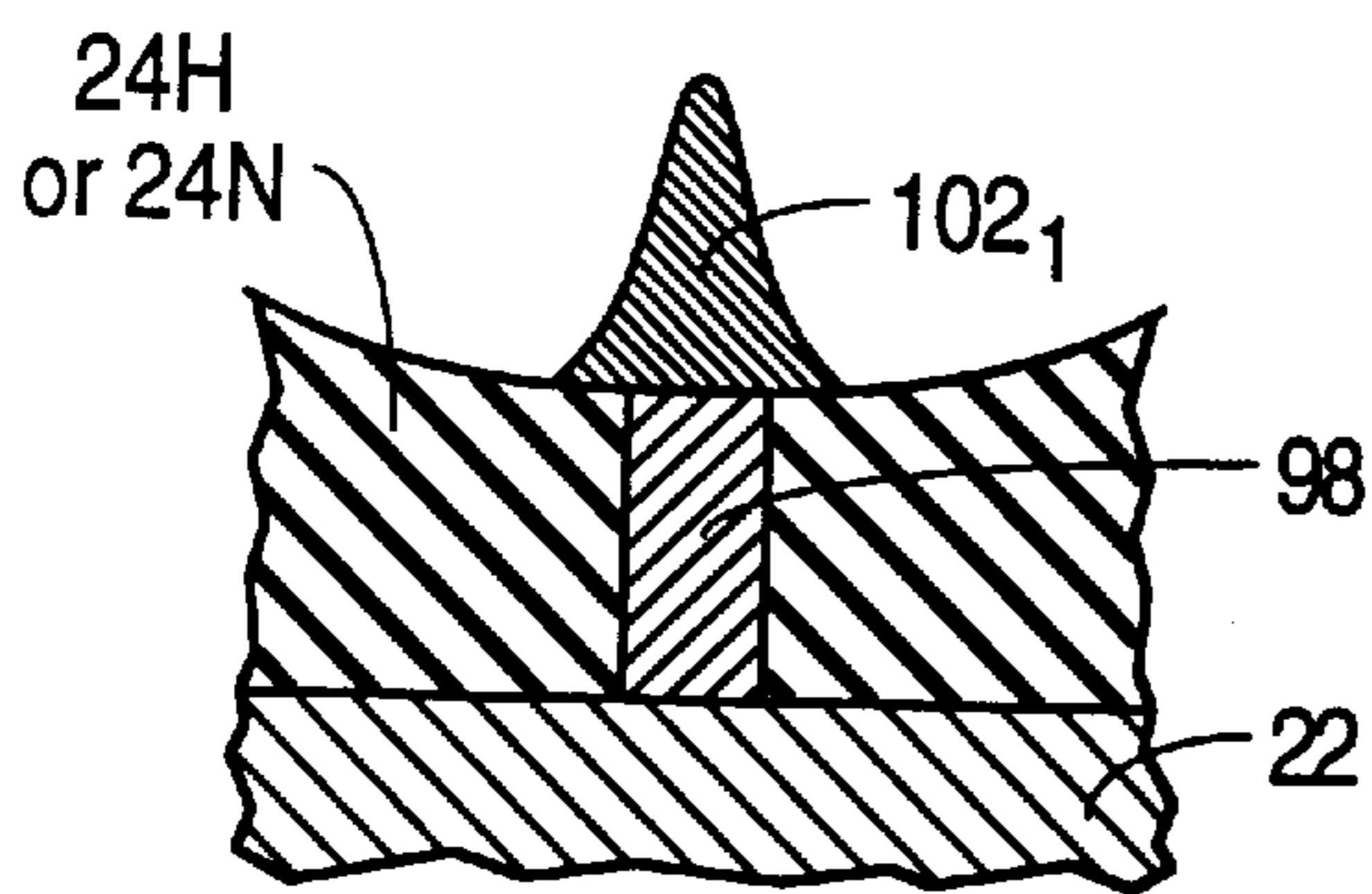


FIG. 16.1

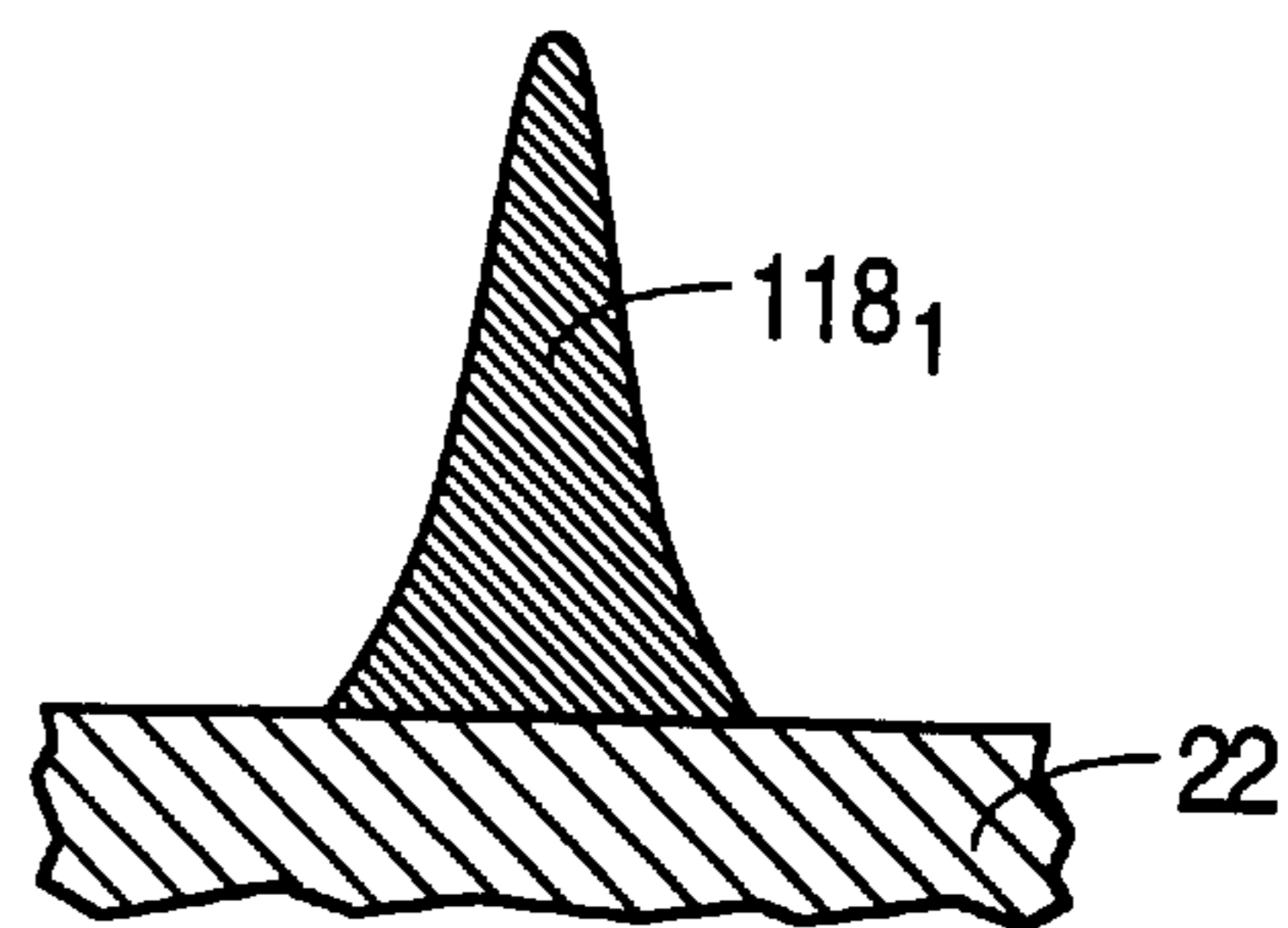


FIG. 16.2

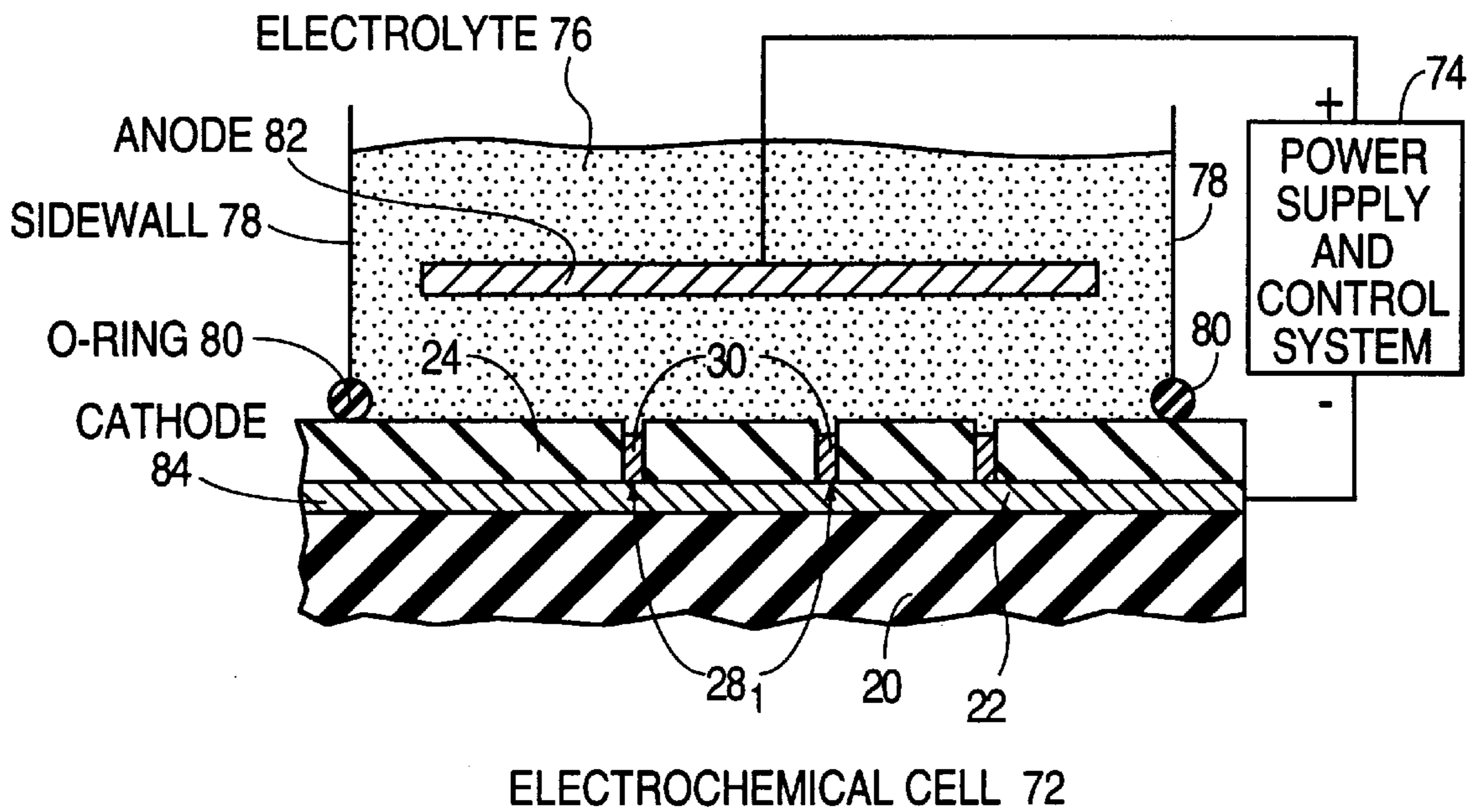
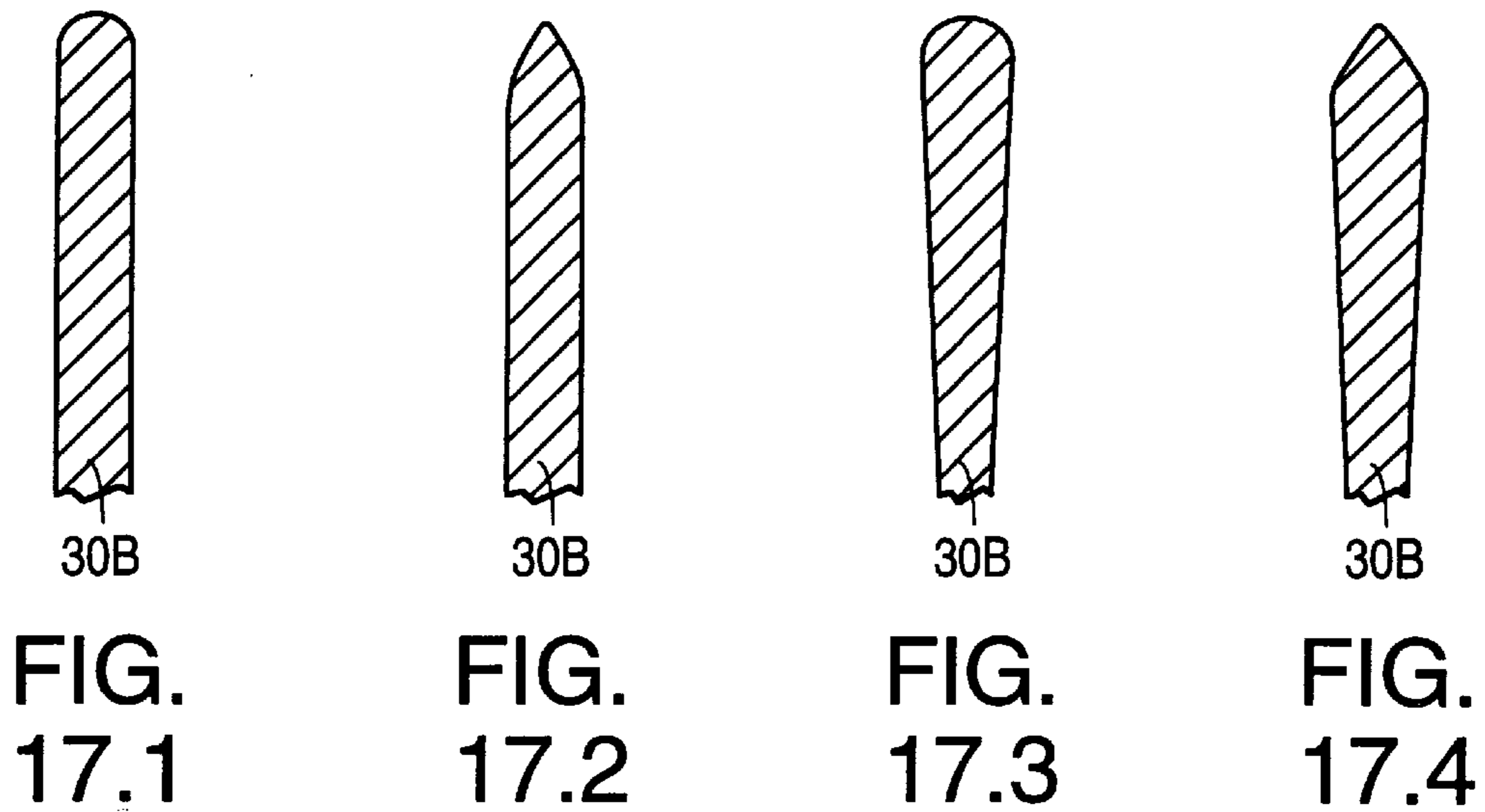


FIG. 18

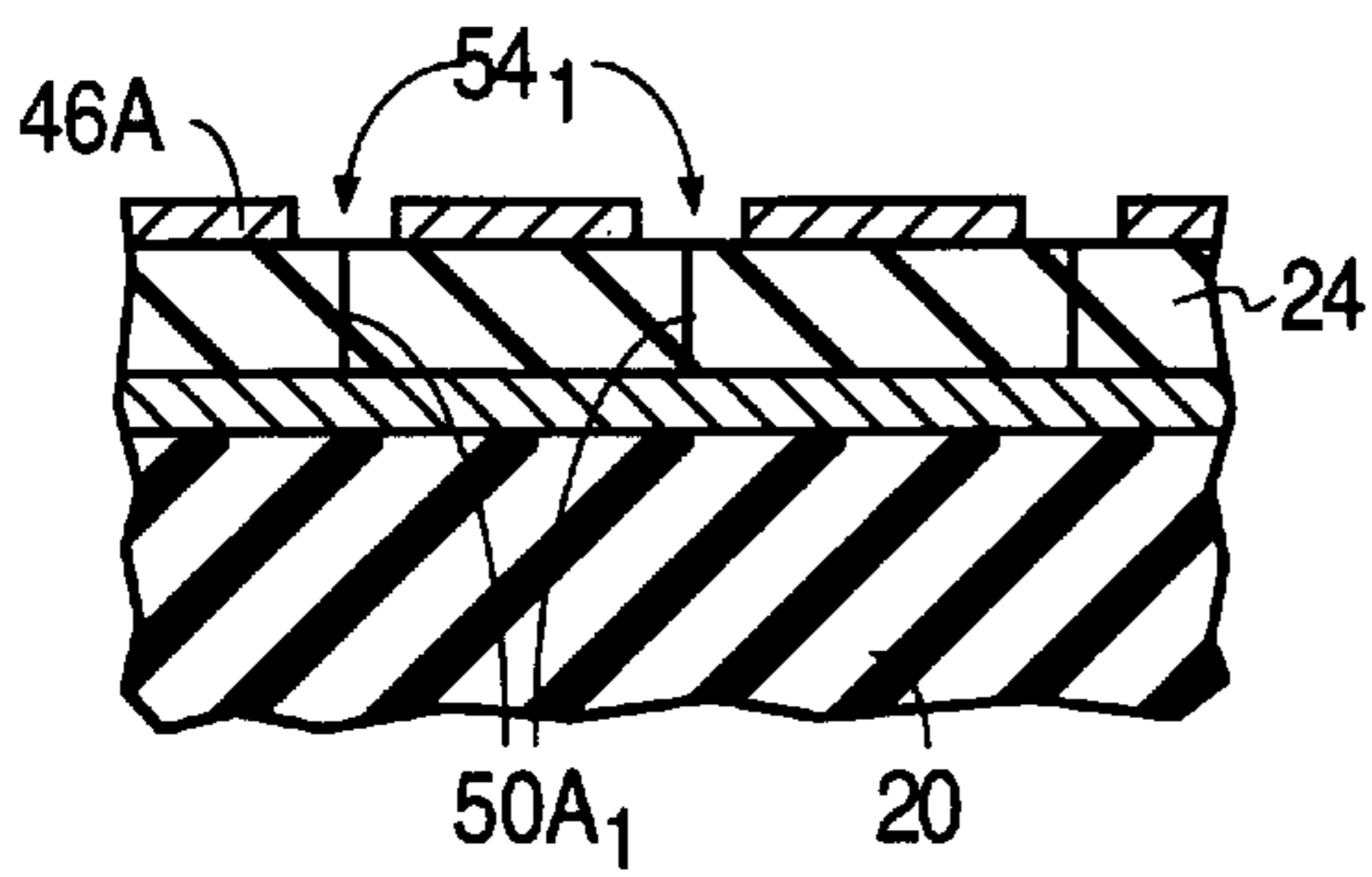


FIG. 19a

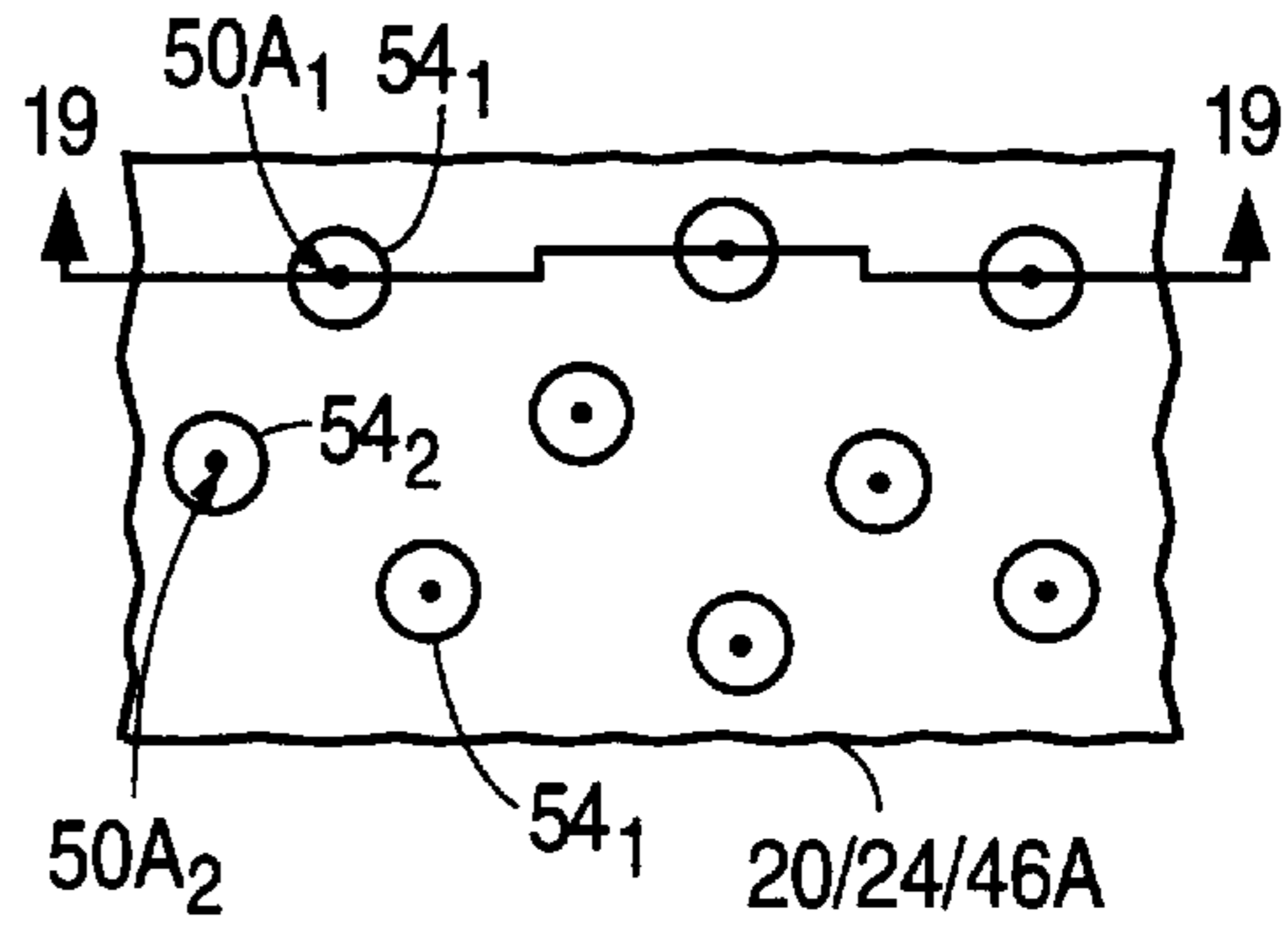


FIG. 20a

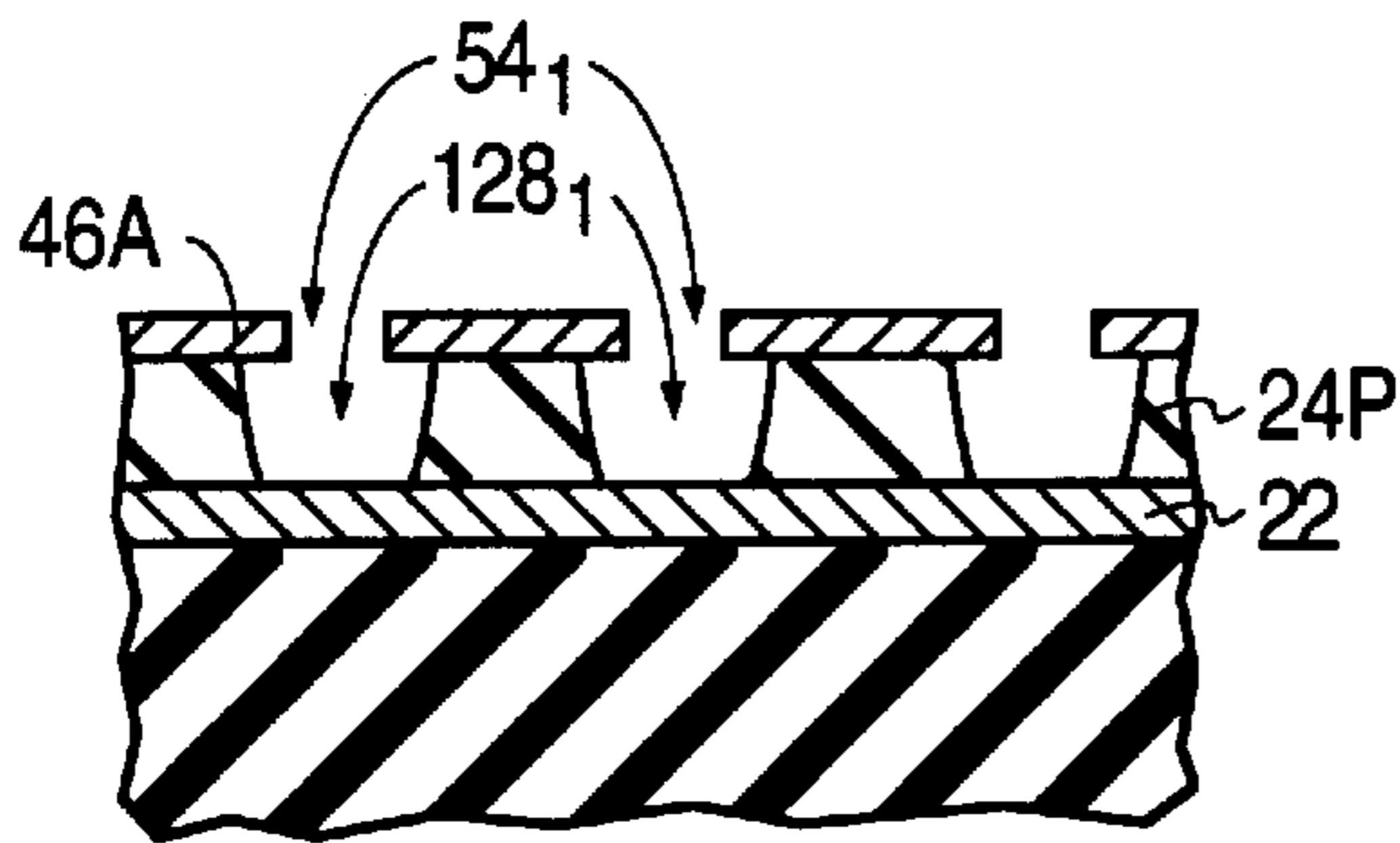


FIG. 19b

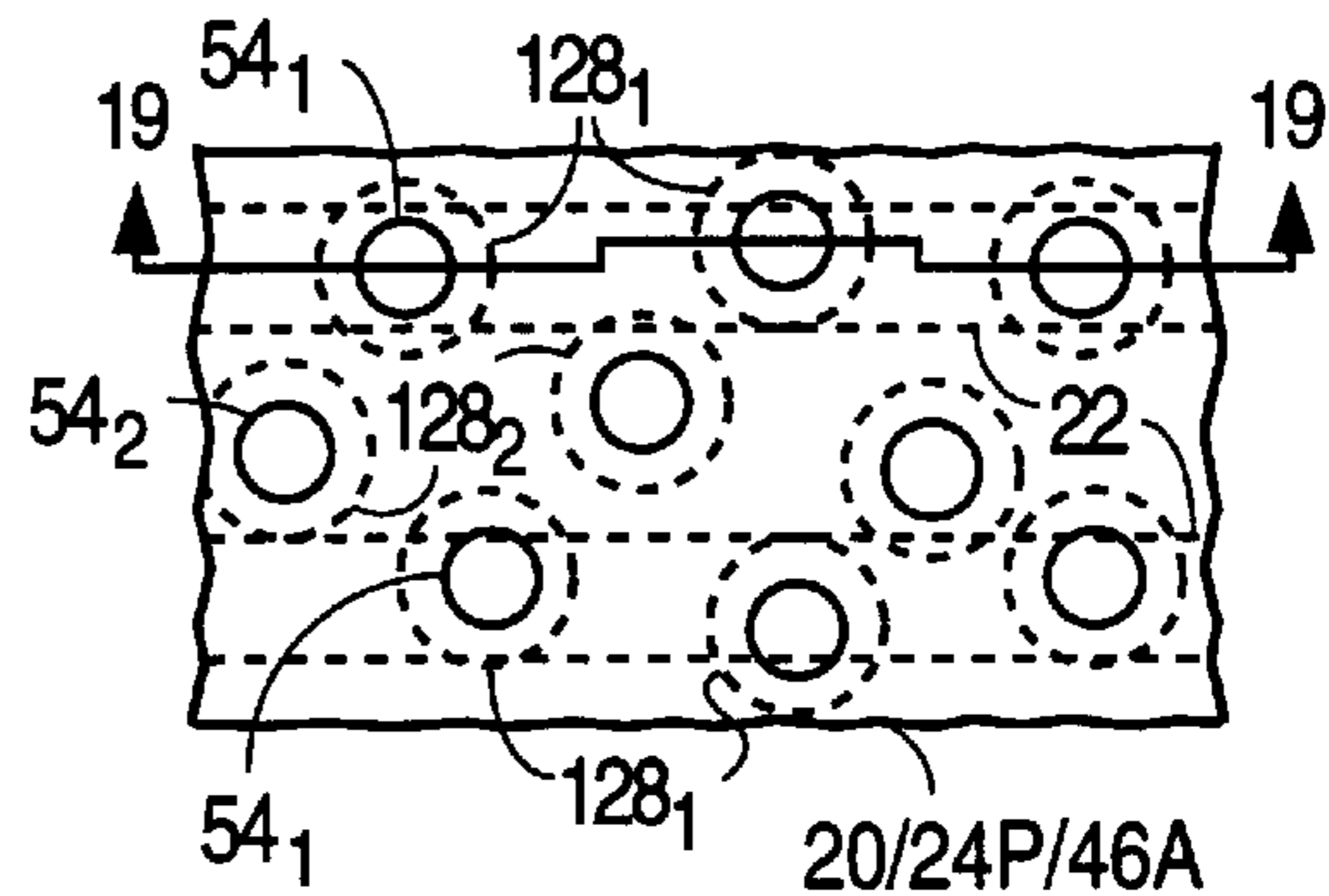


FIG. 20b

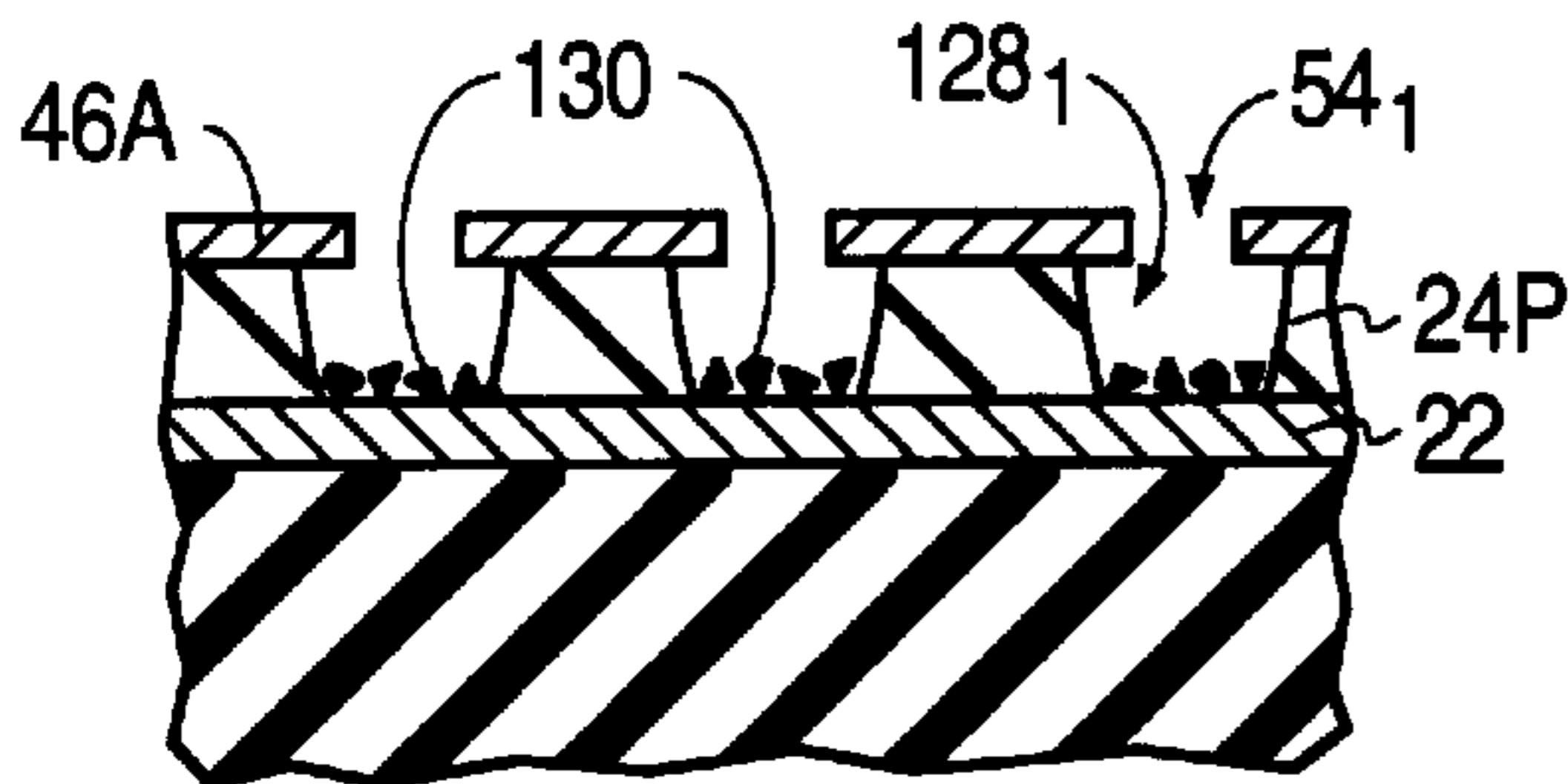


FIG. 19c

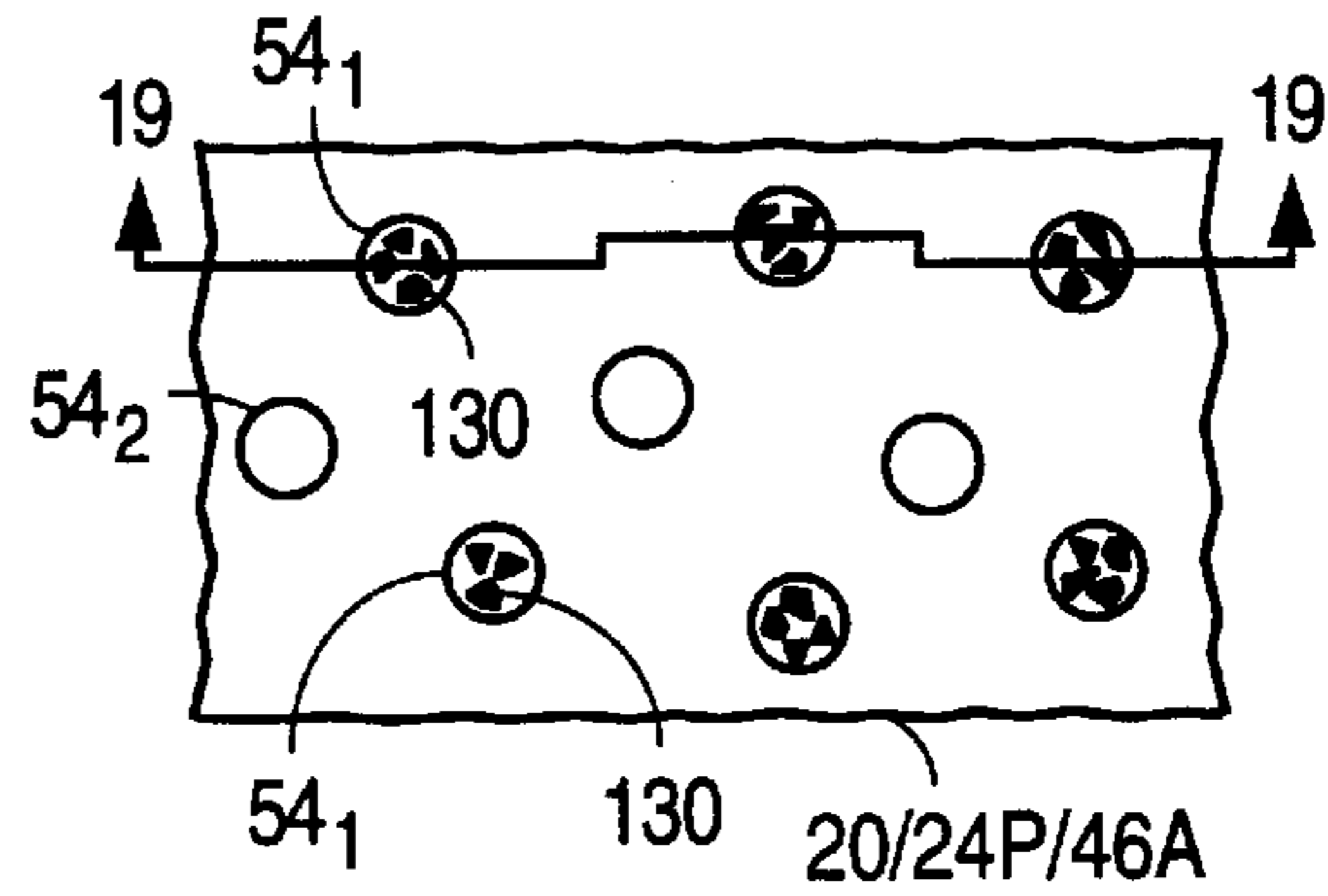


FIG. 20c

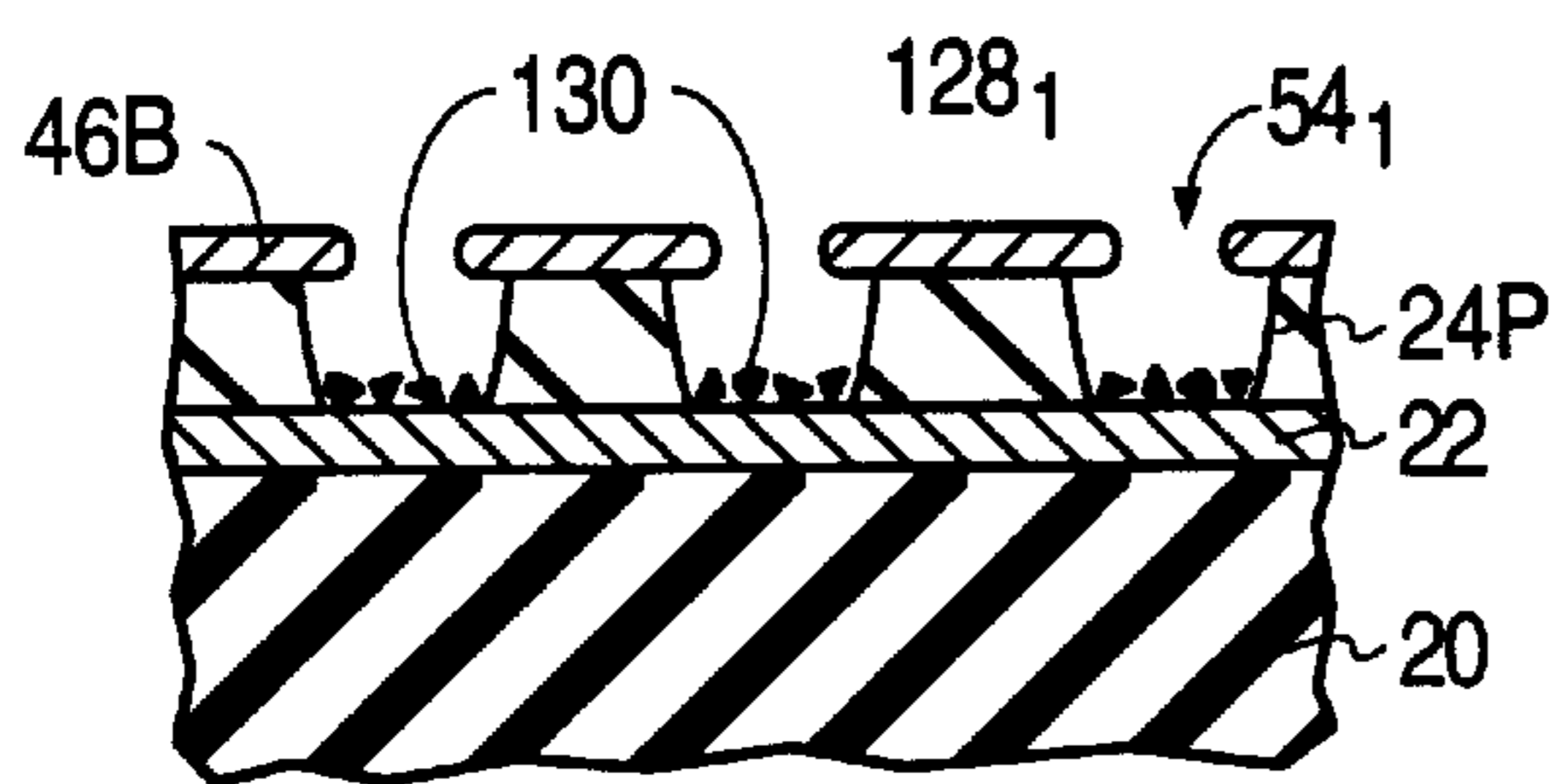


FIG. 19d

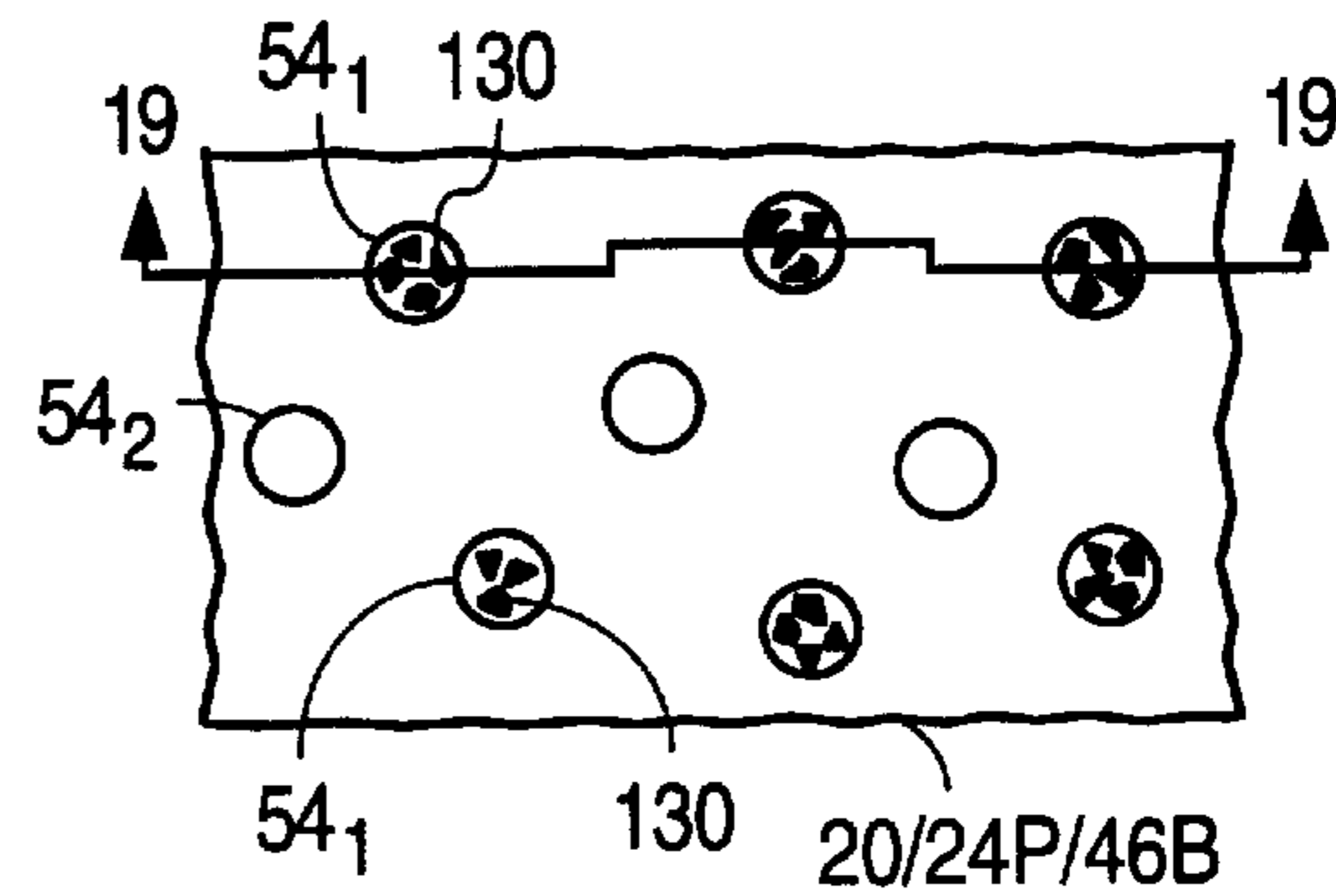


FIG. 20d

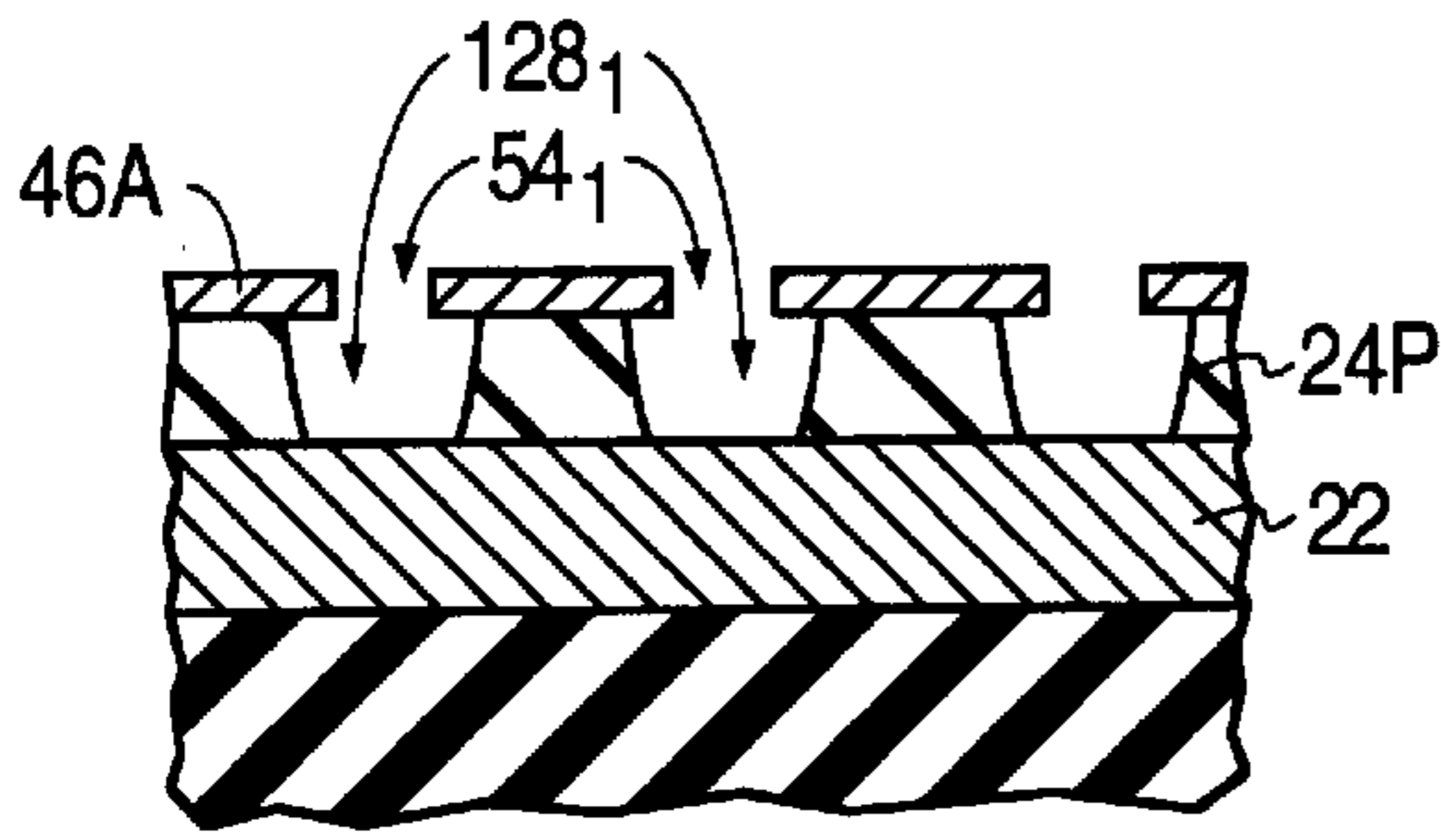
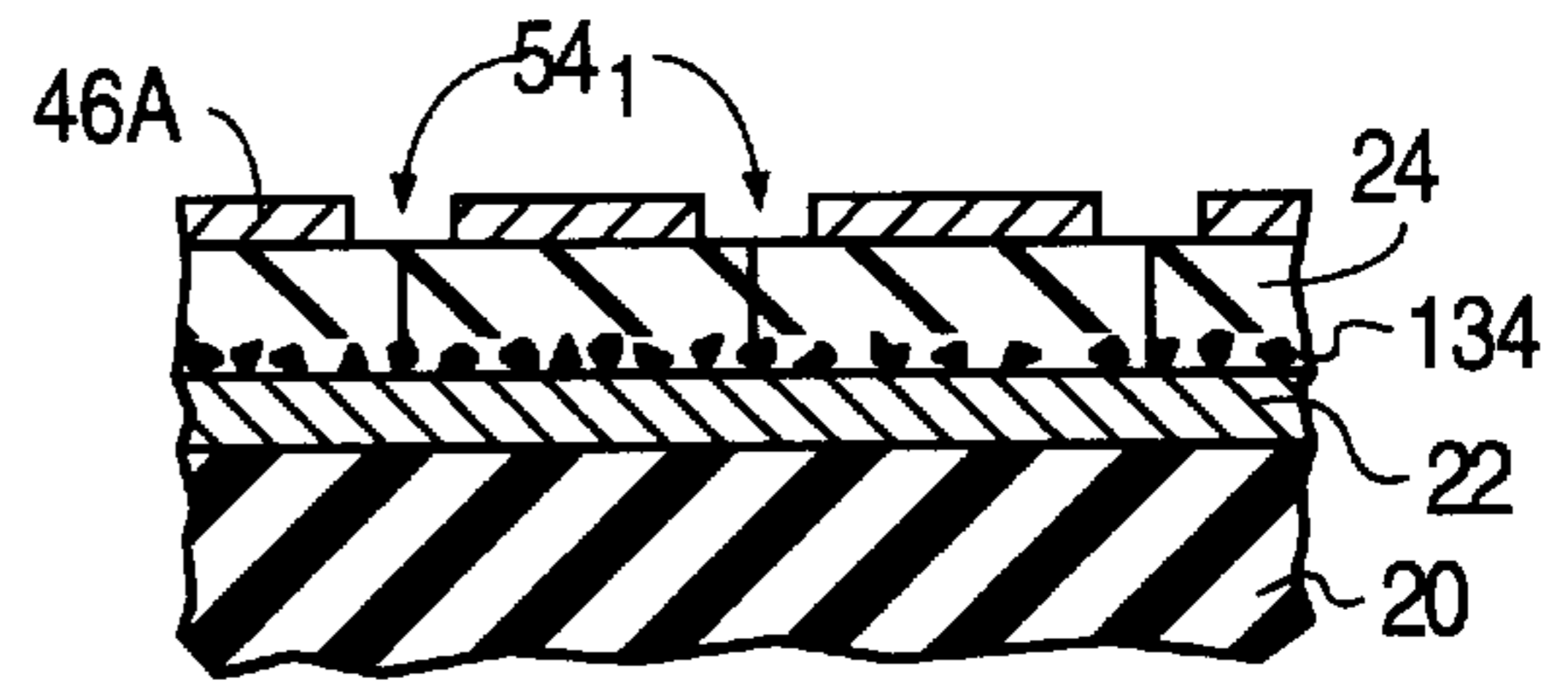
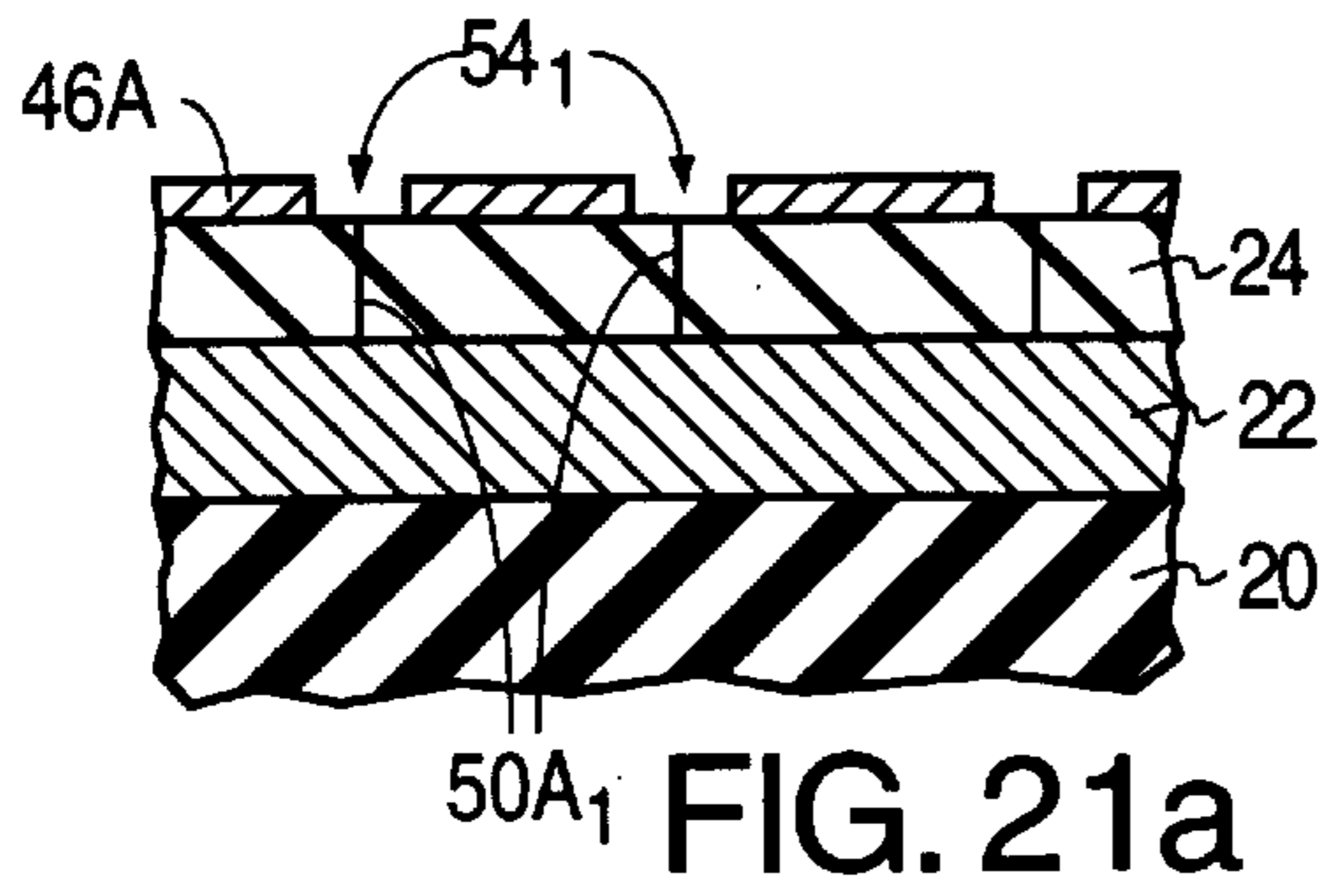


FIG. 22a

FIG. 21b

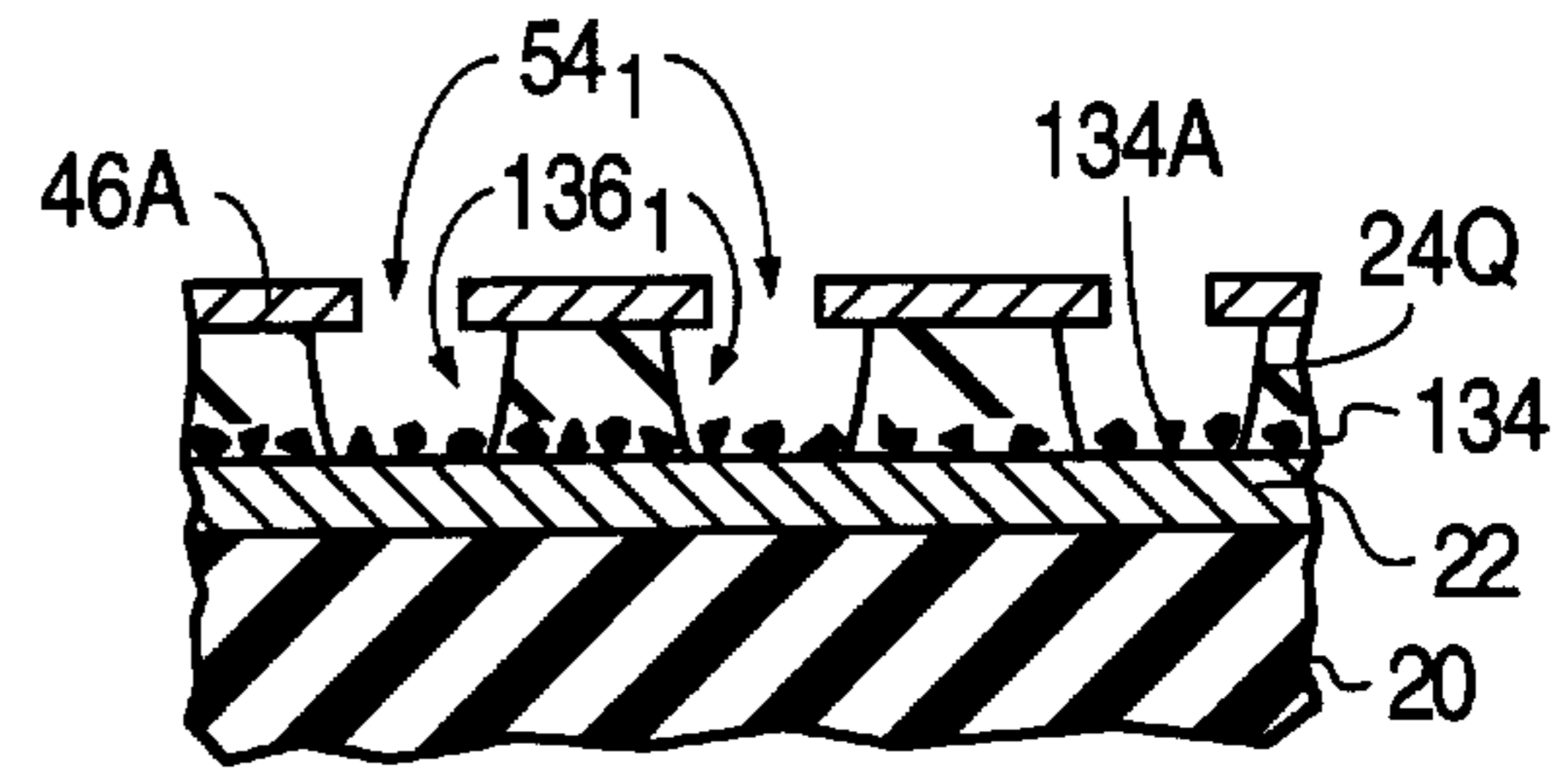
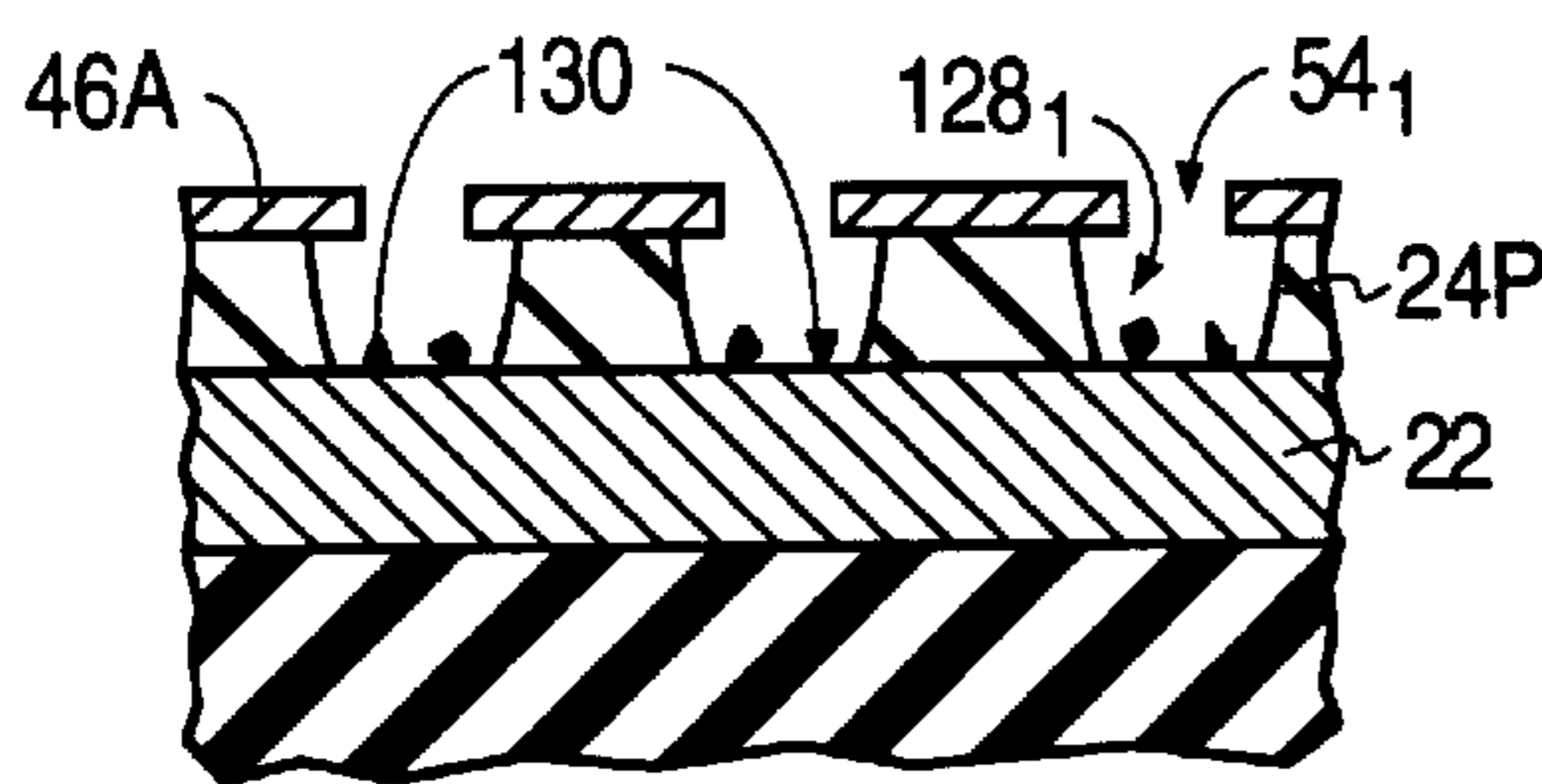


FIG. 22b

FIG. 21c

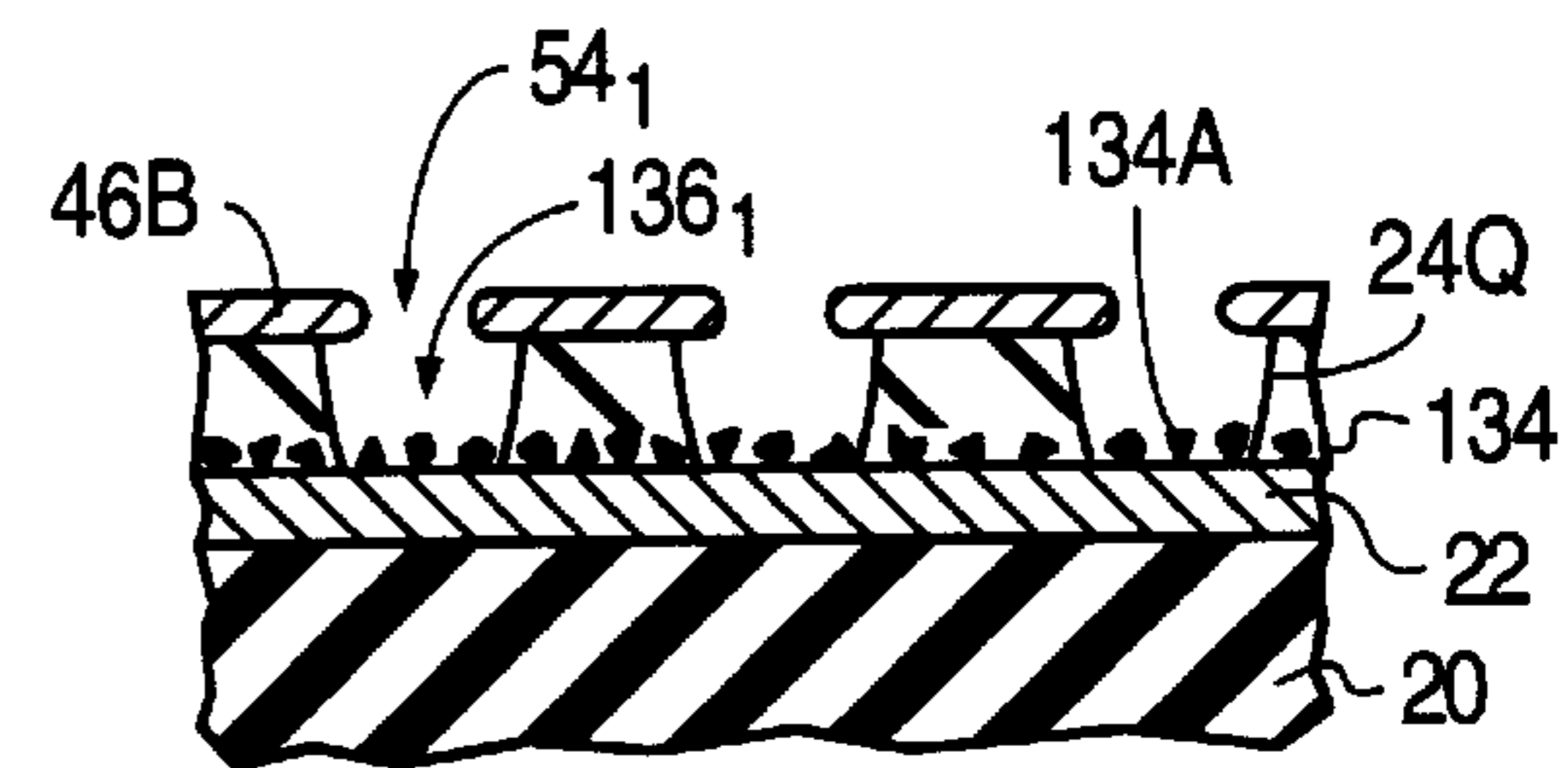
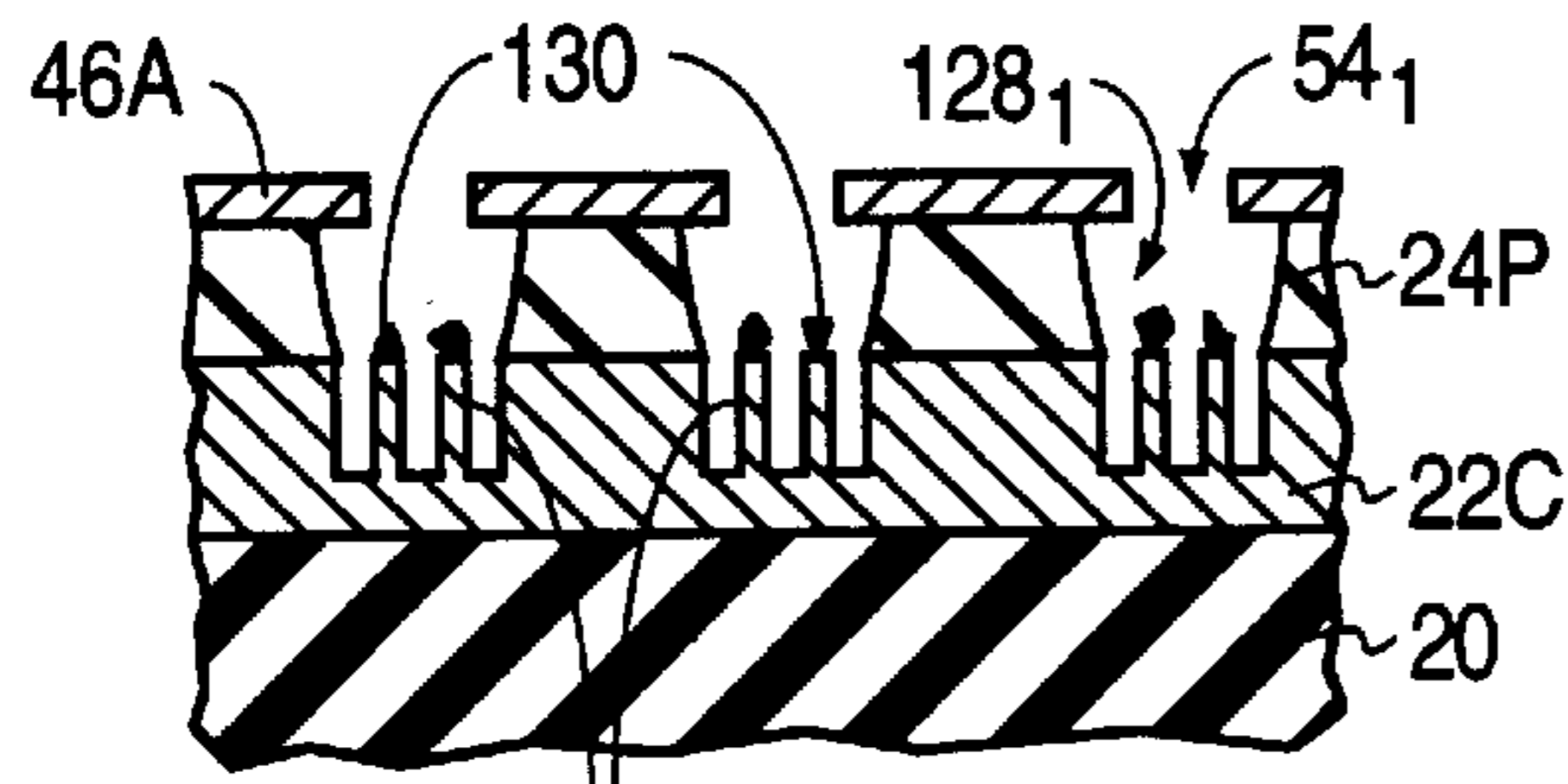


FIG. 22c

FIG. 21d

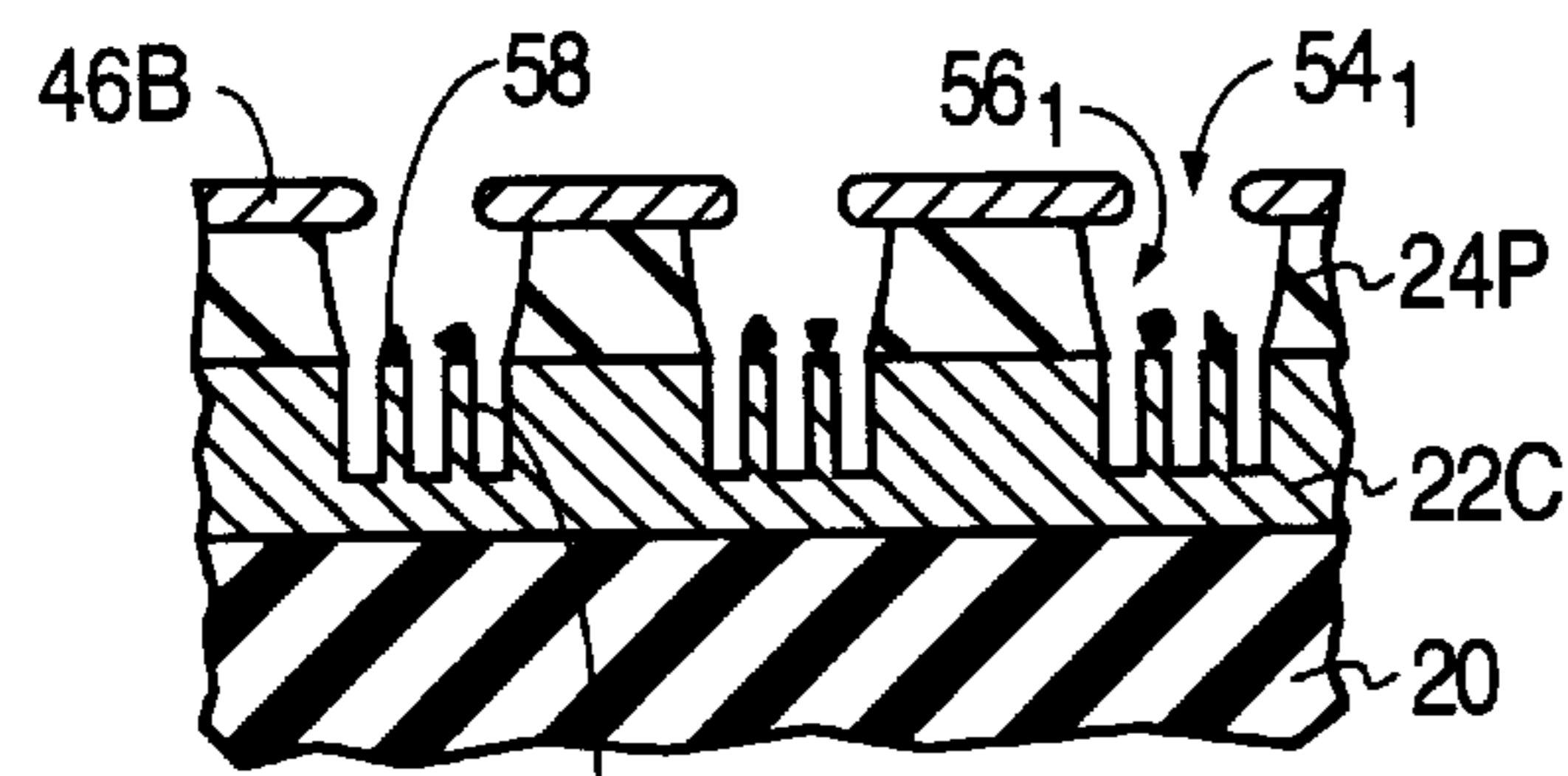
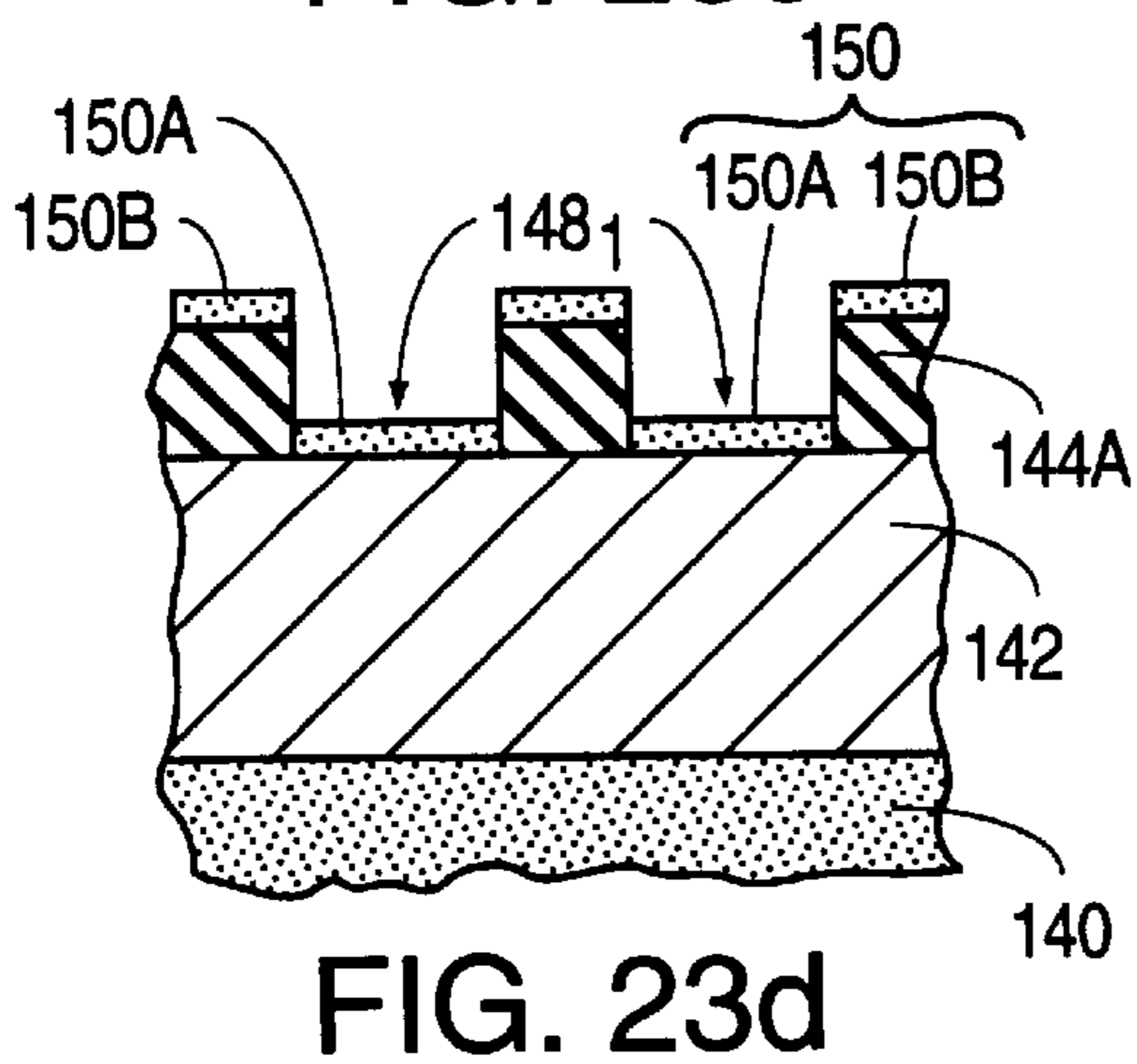
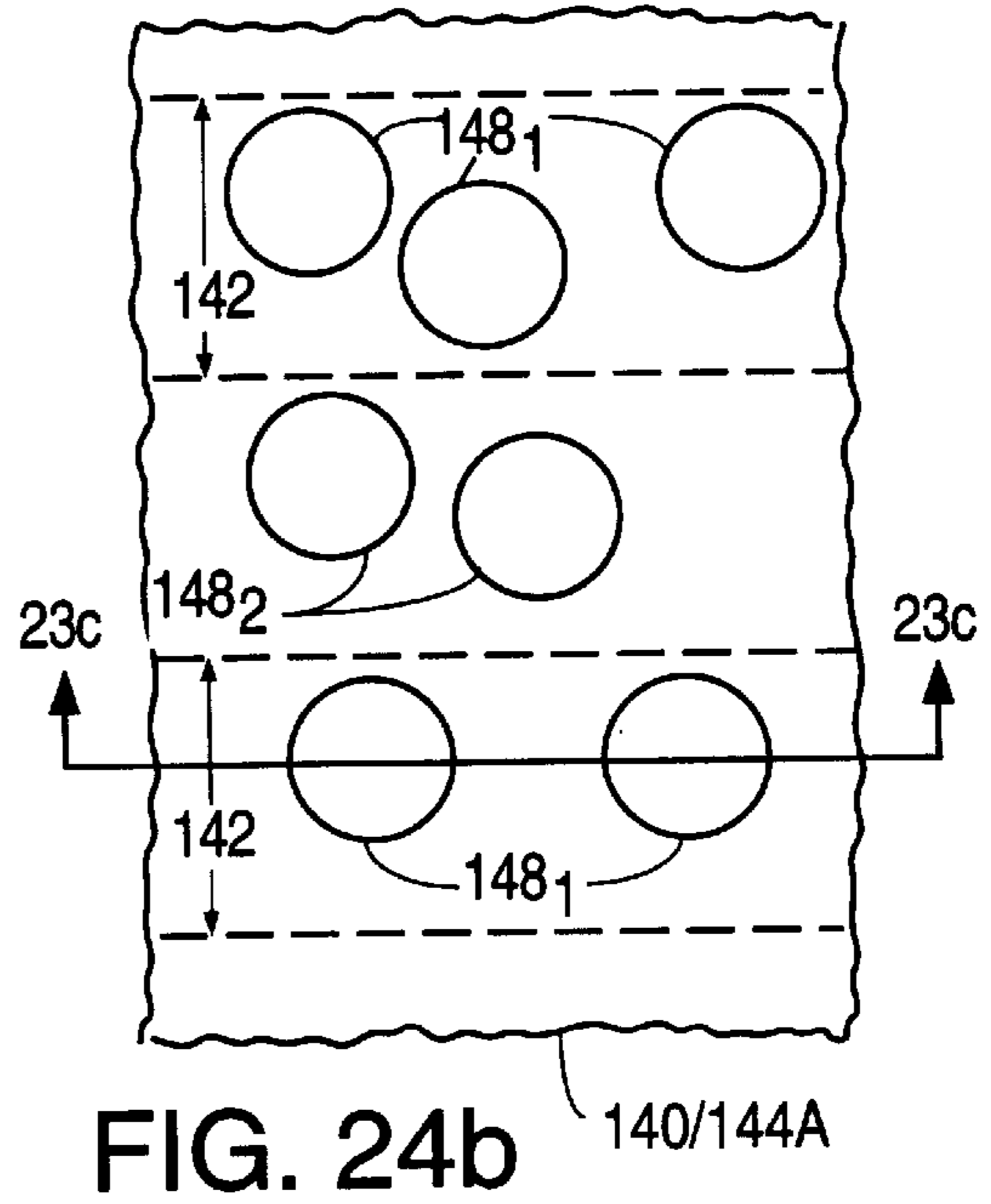
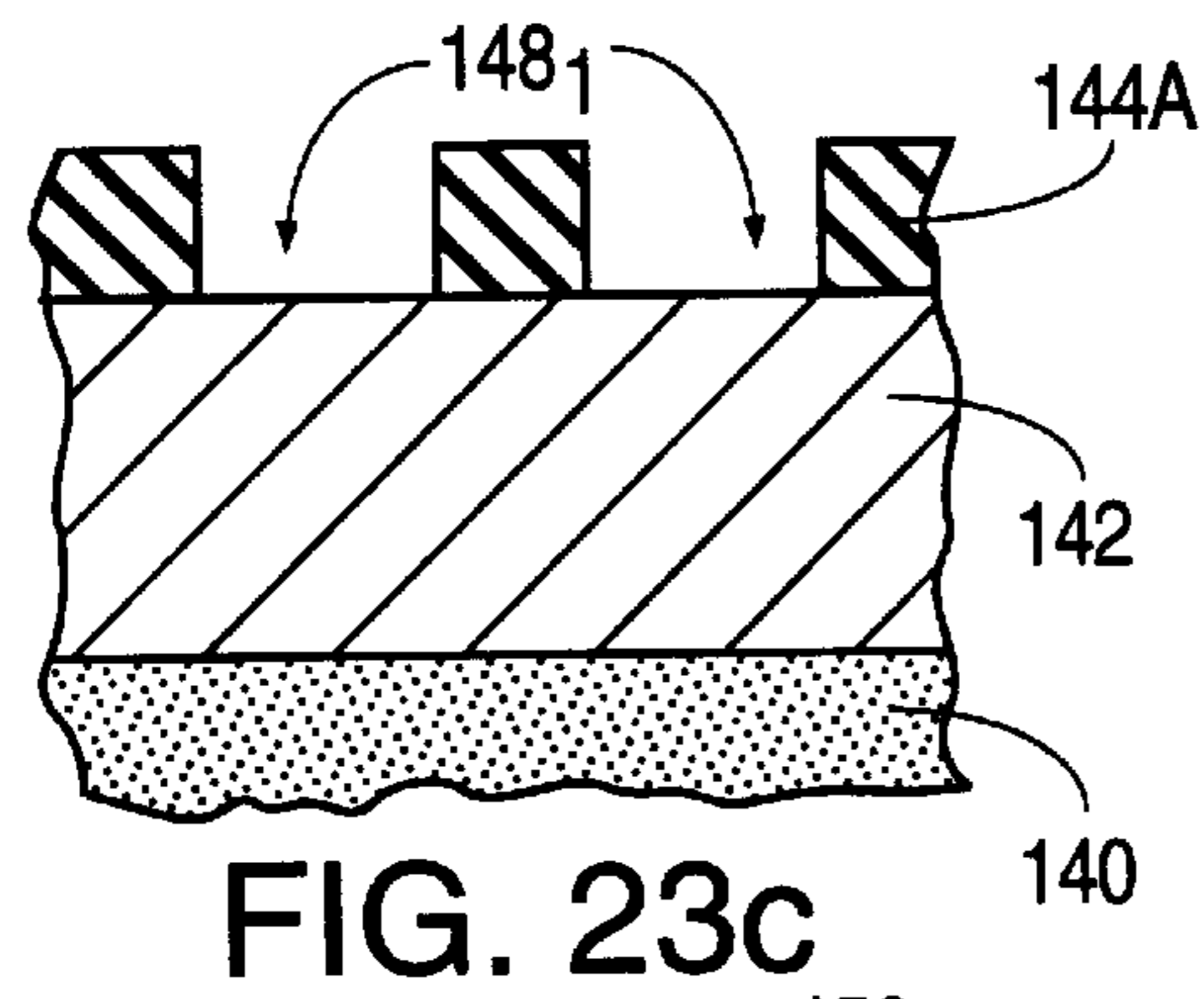
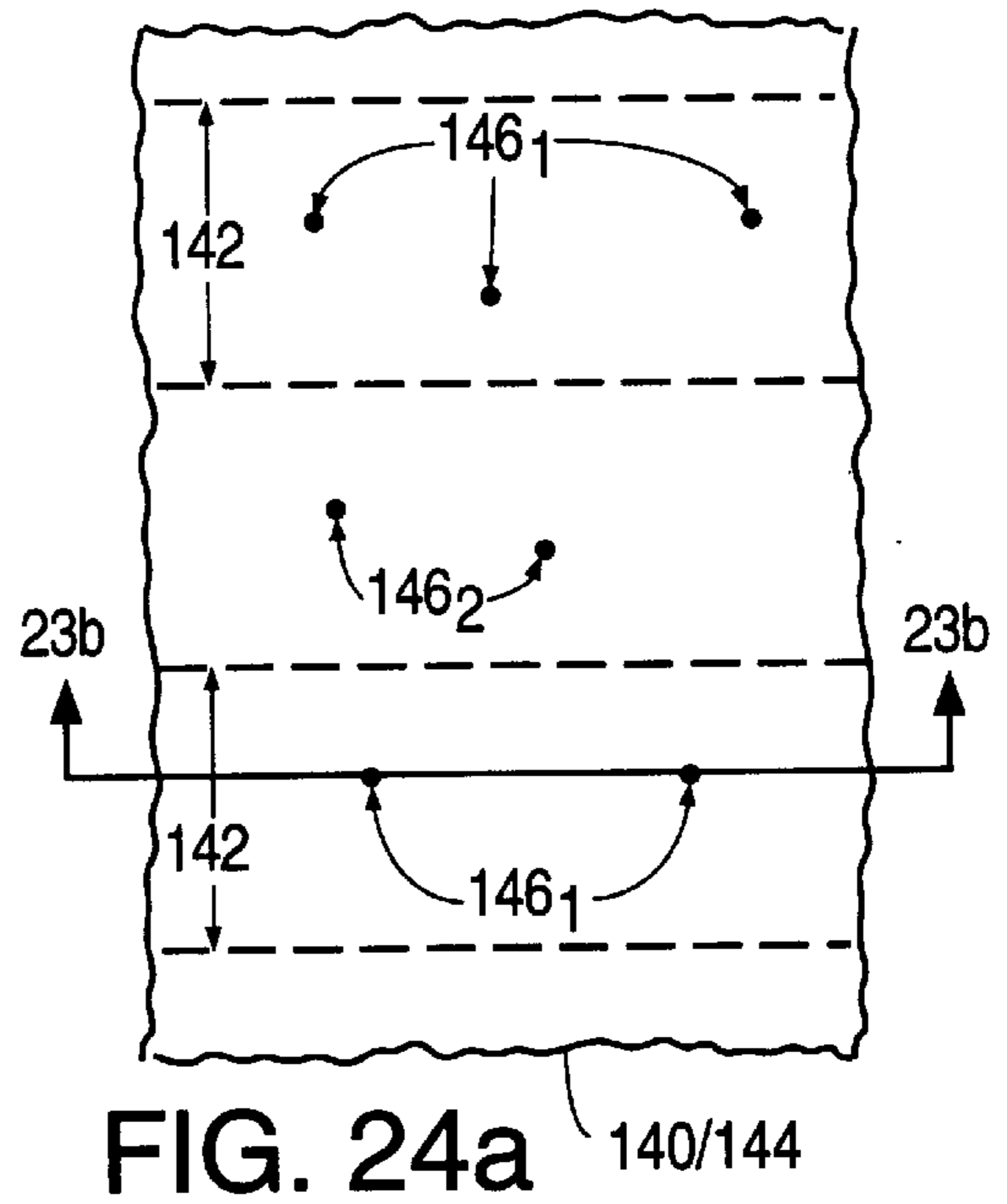
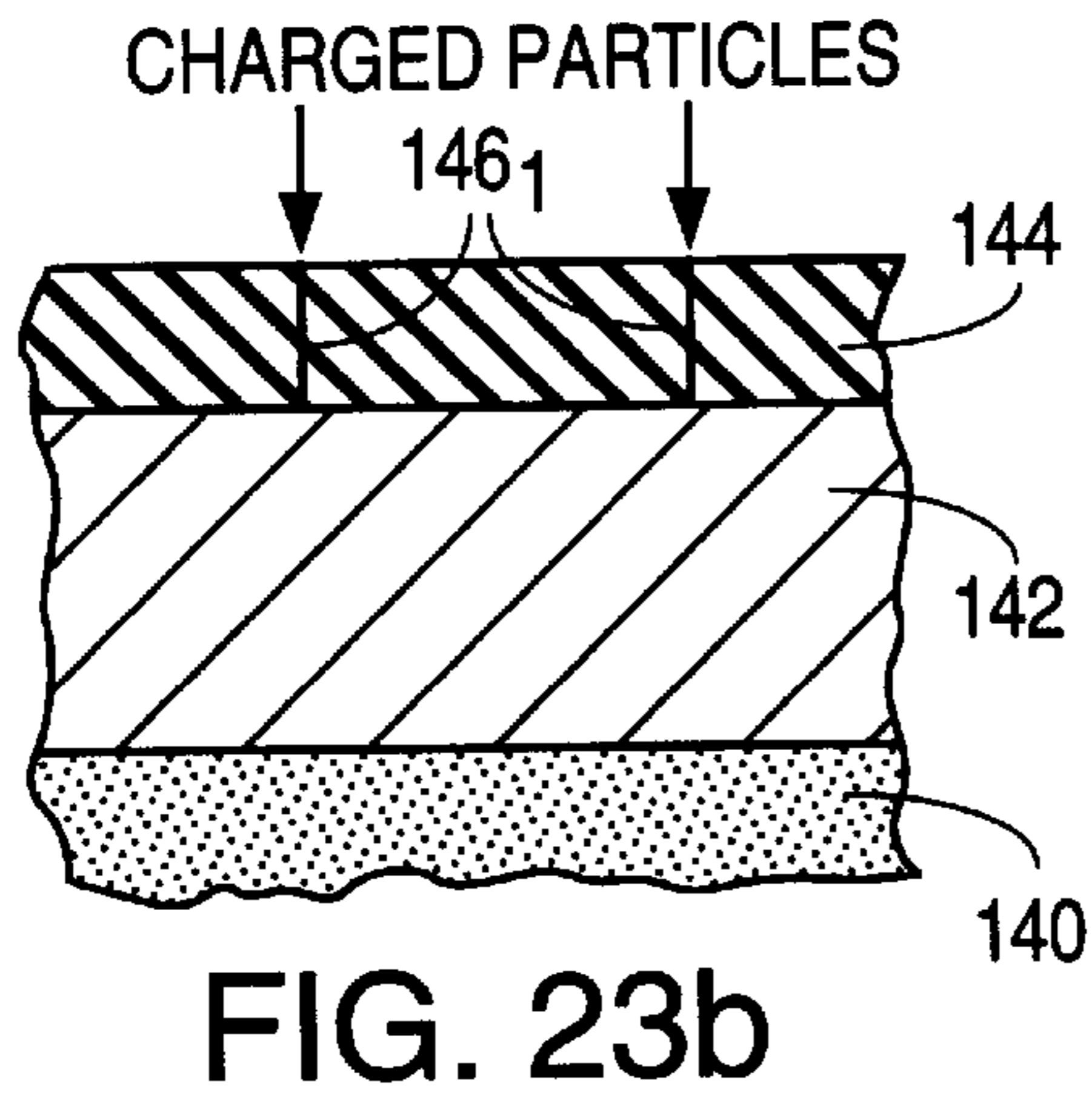
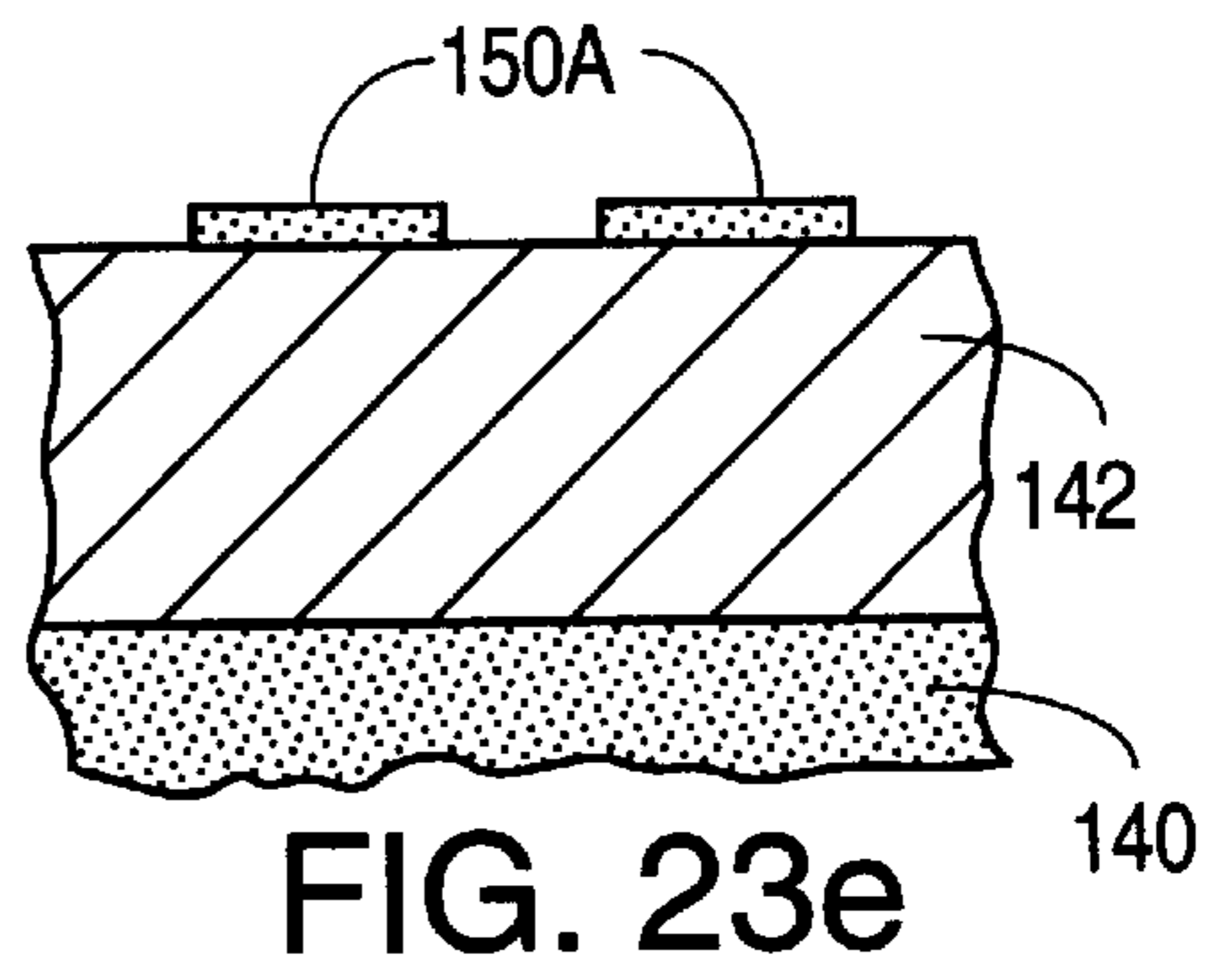
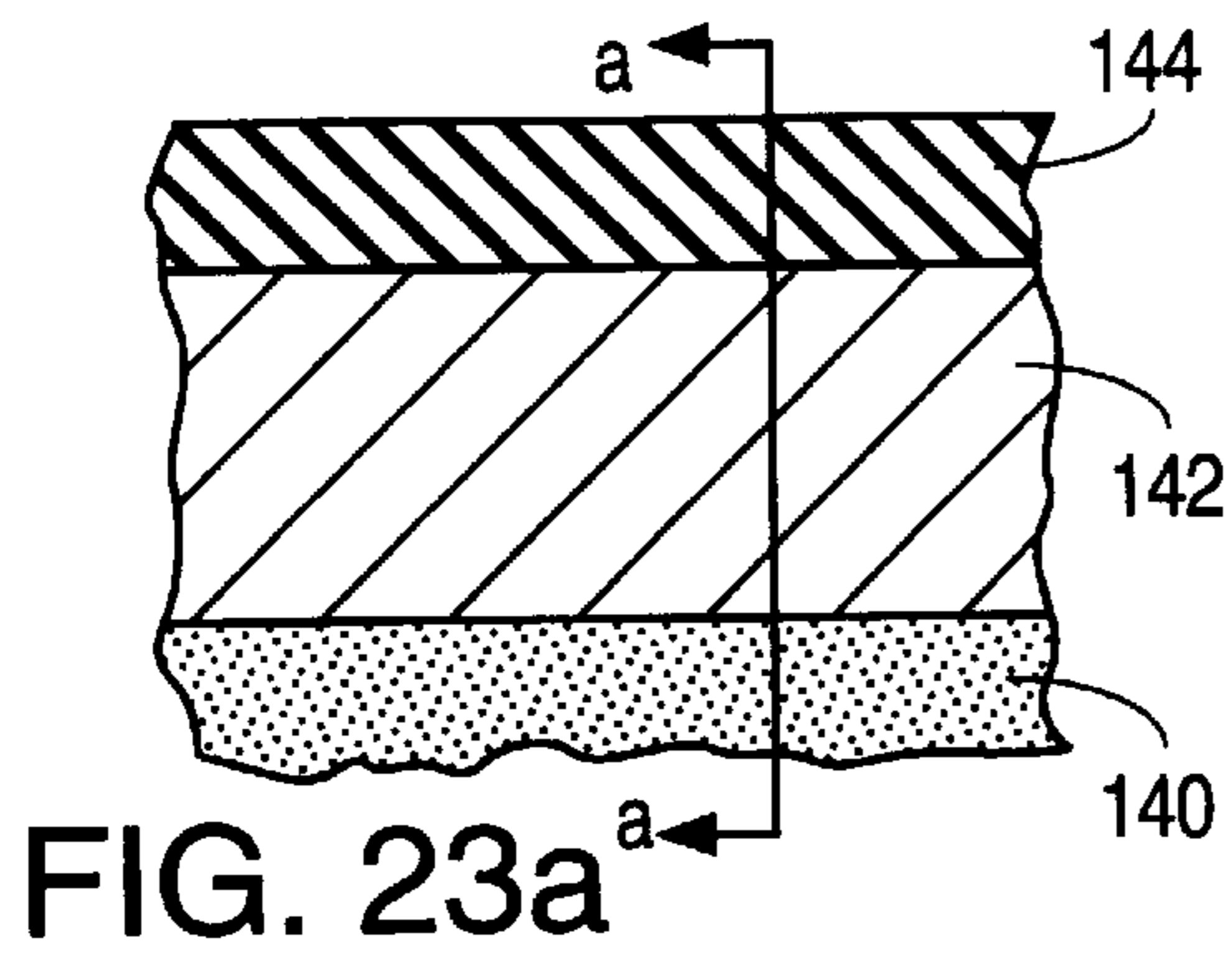


FIG. 21e



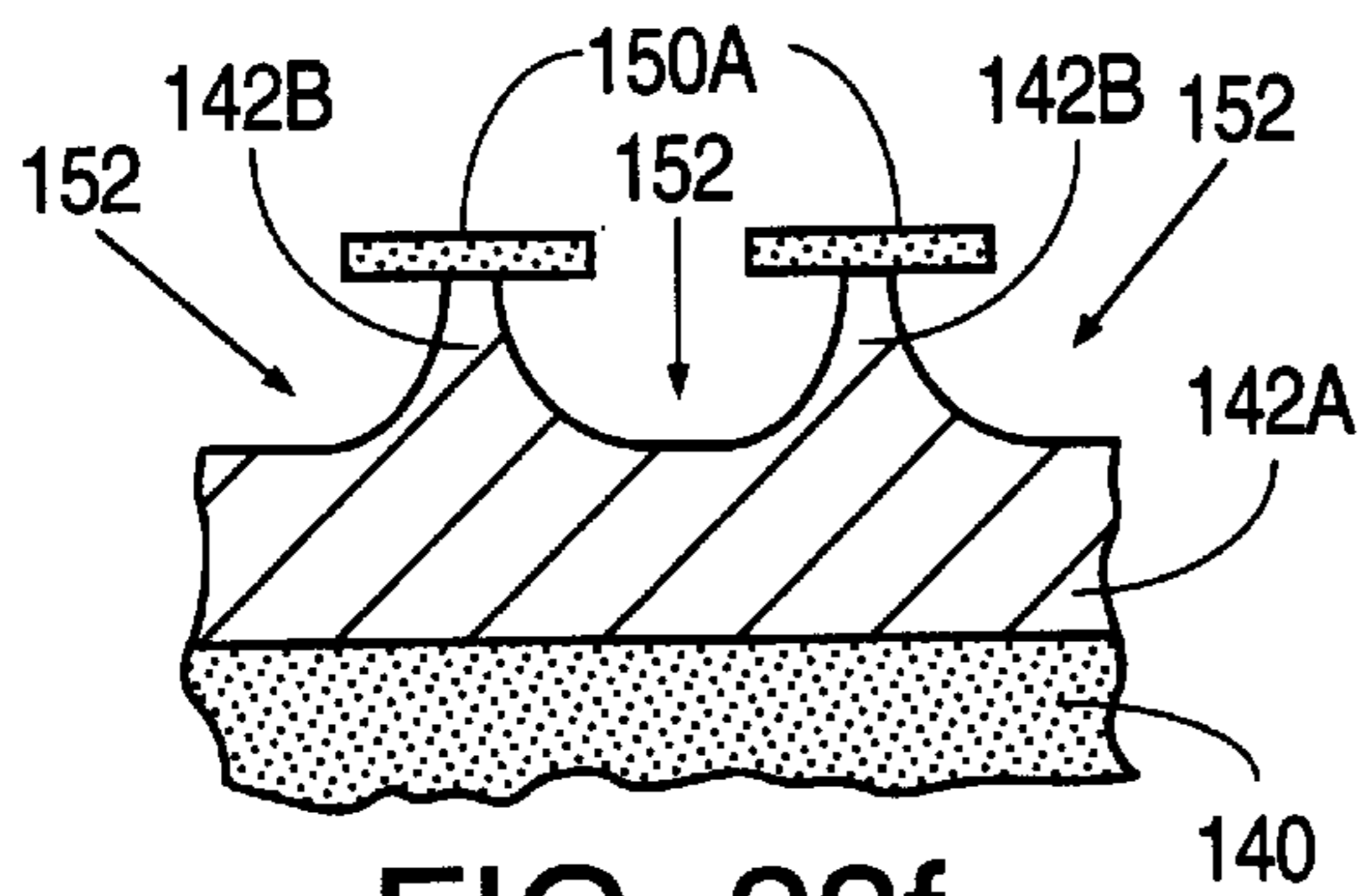


FIG. 23f

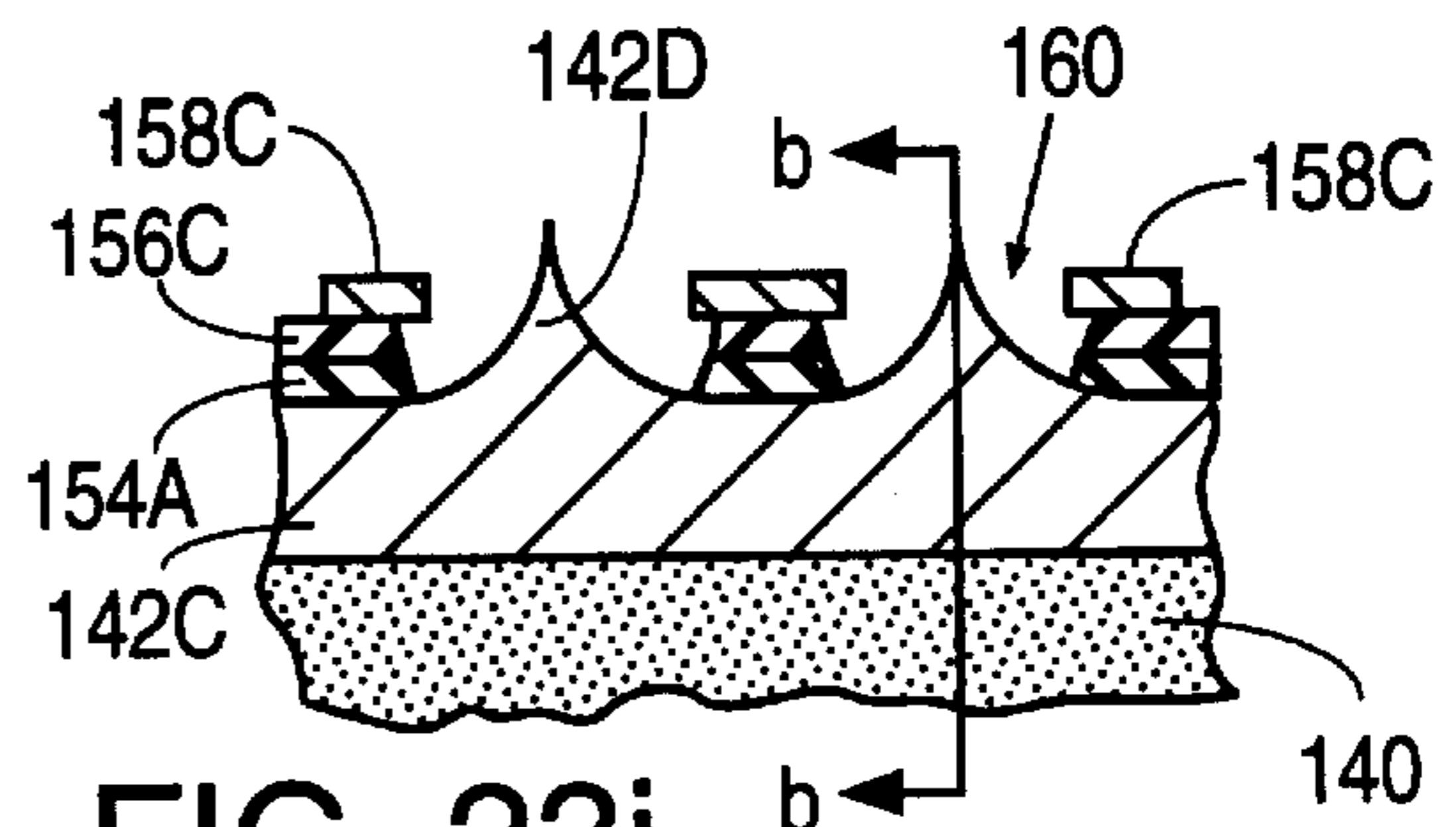


FIG. 23j

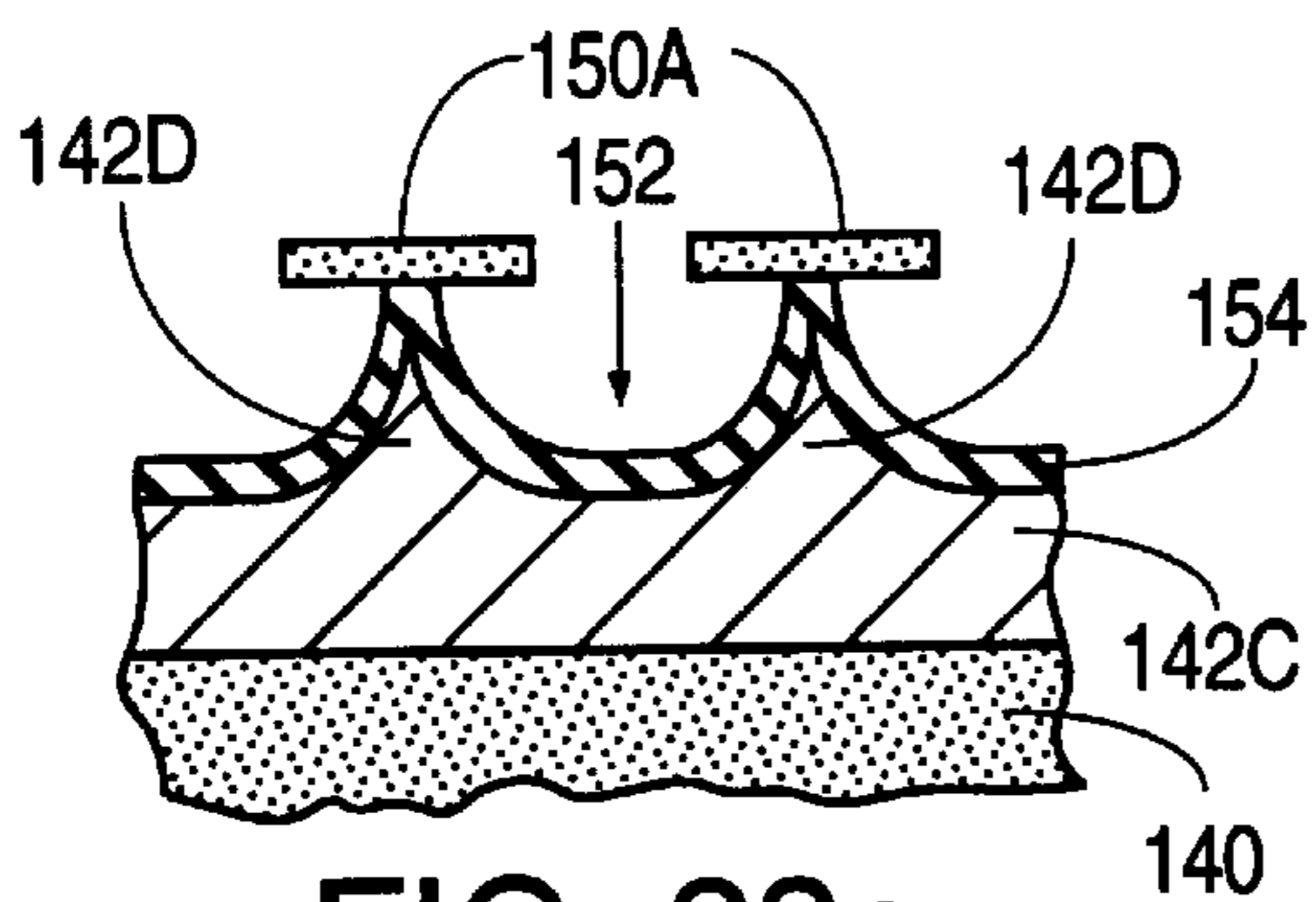


FIG. 23g

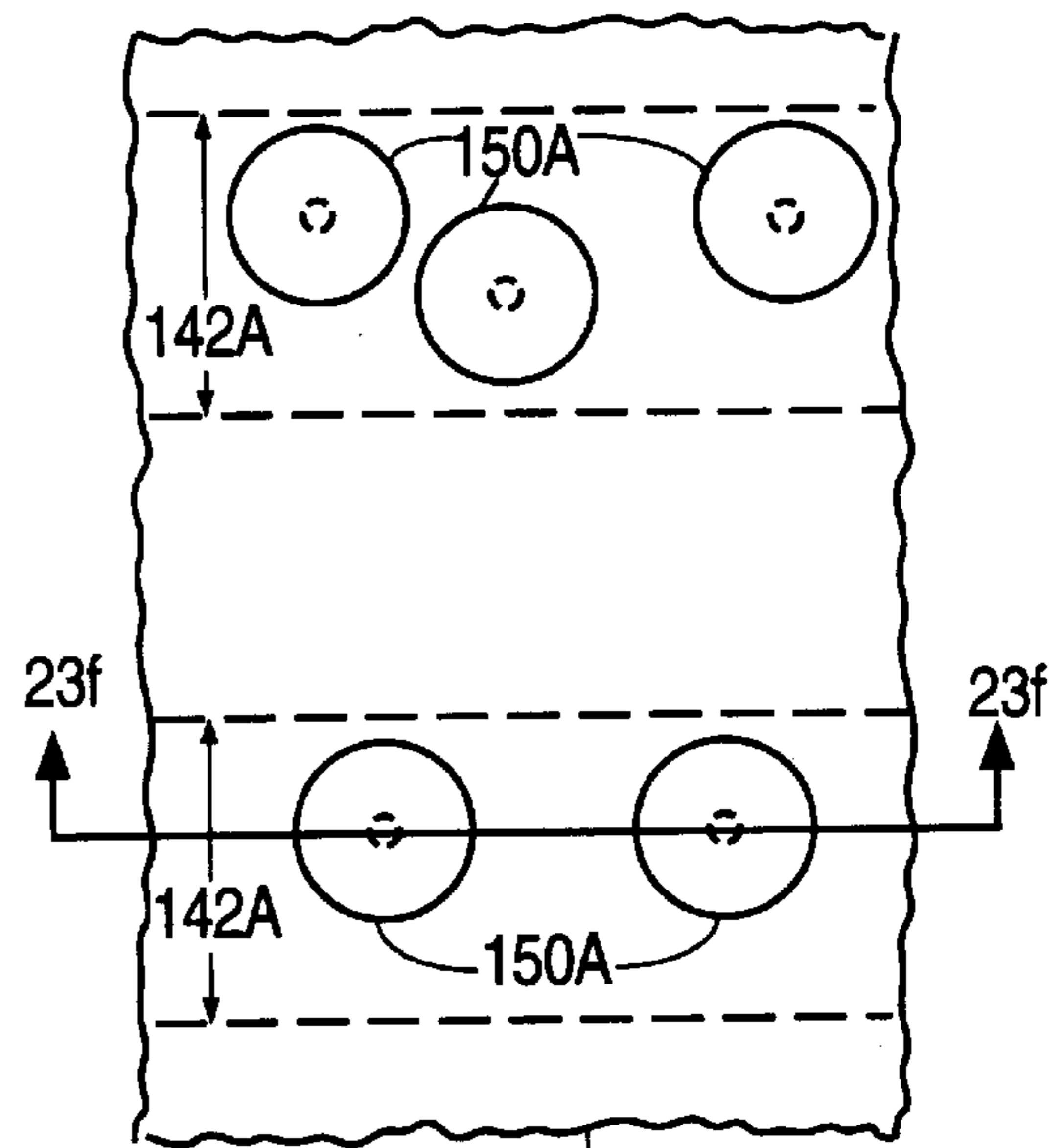


FIG. 24c

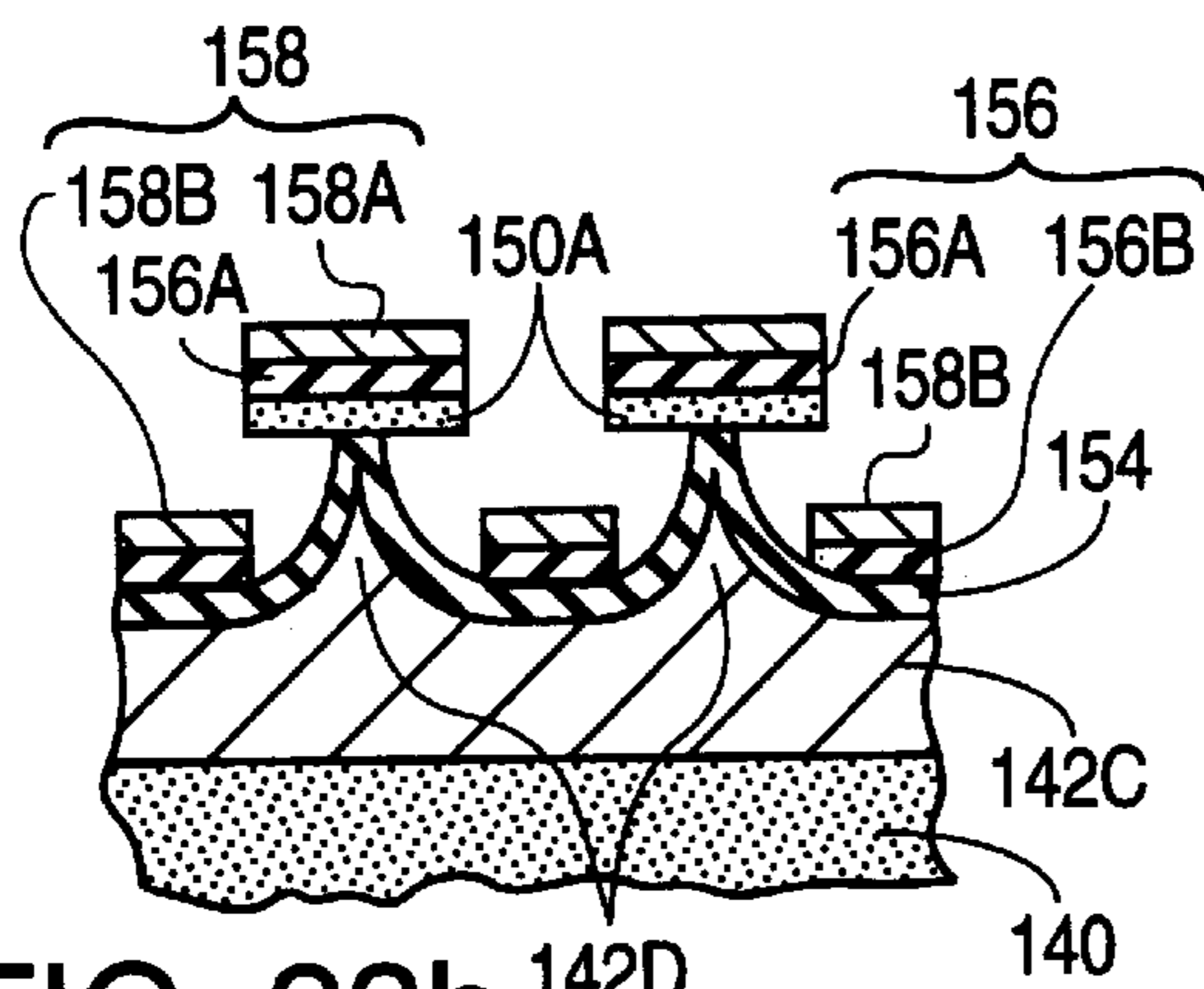


FIG. 23h

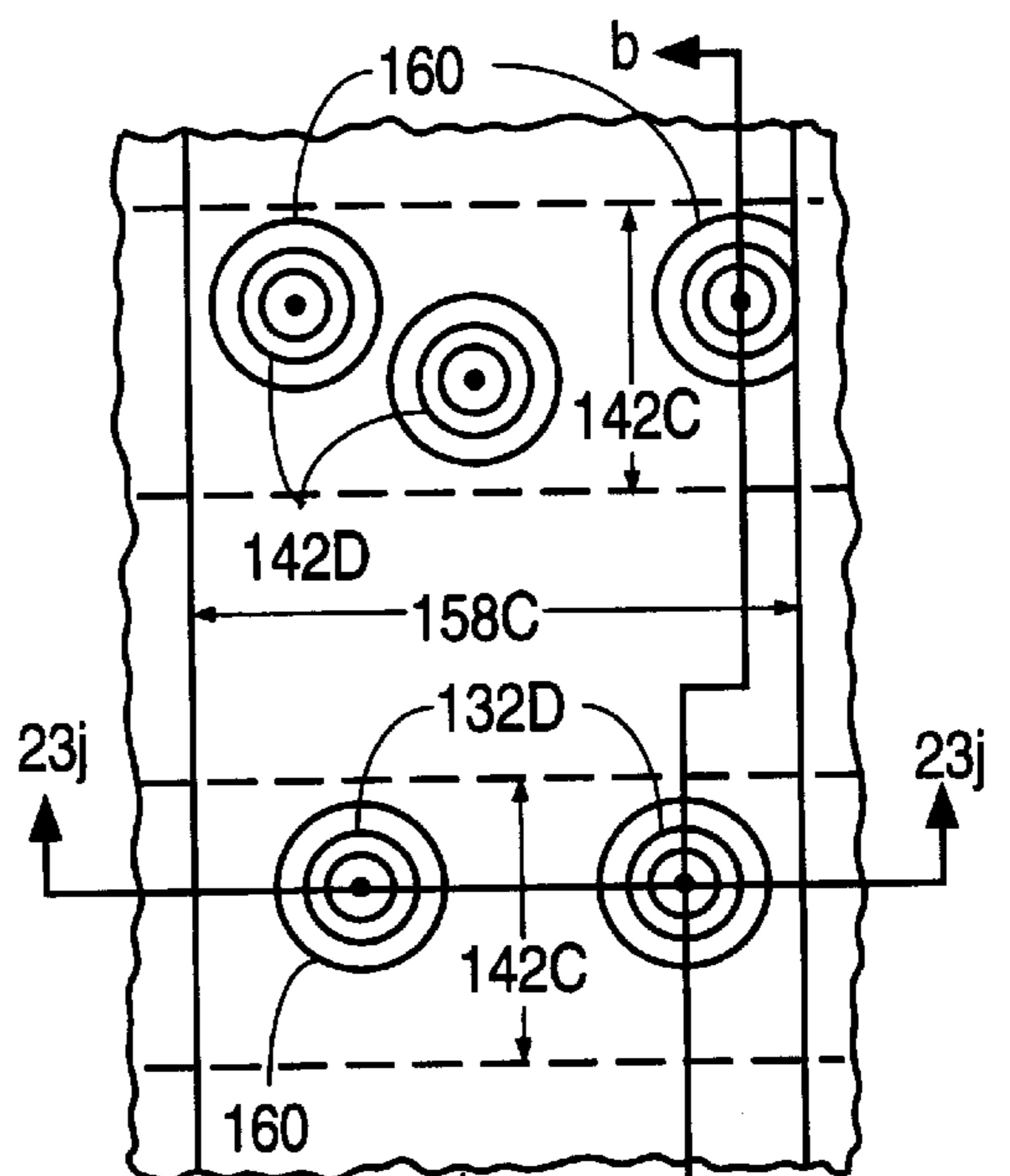


FIG. 24d

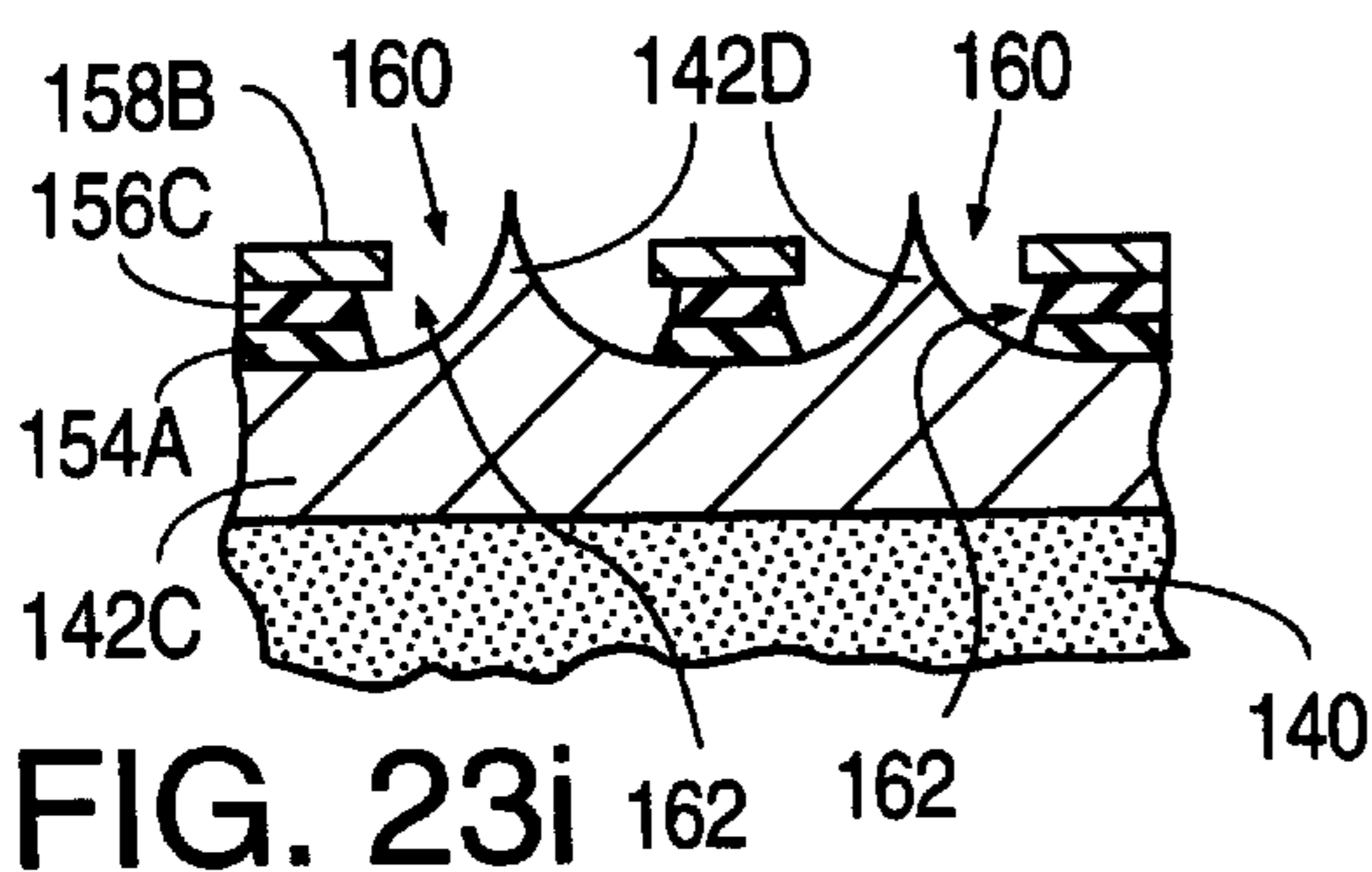


FIG. 23i

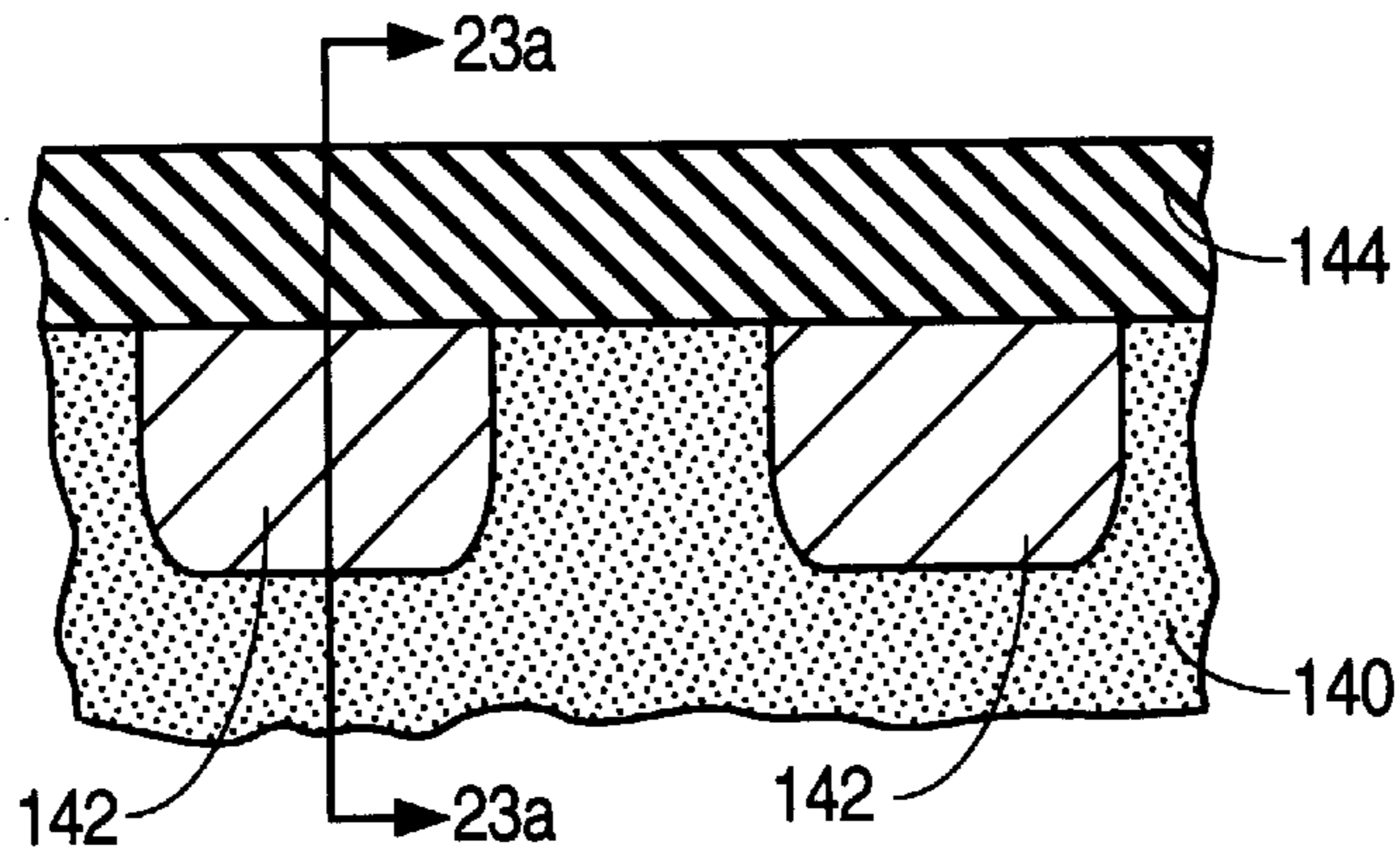


FIG. 25a

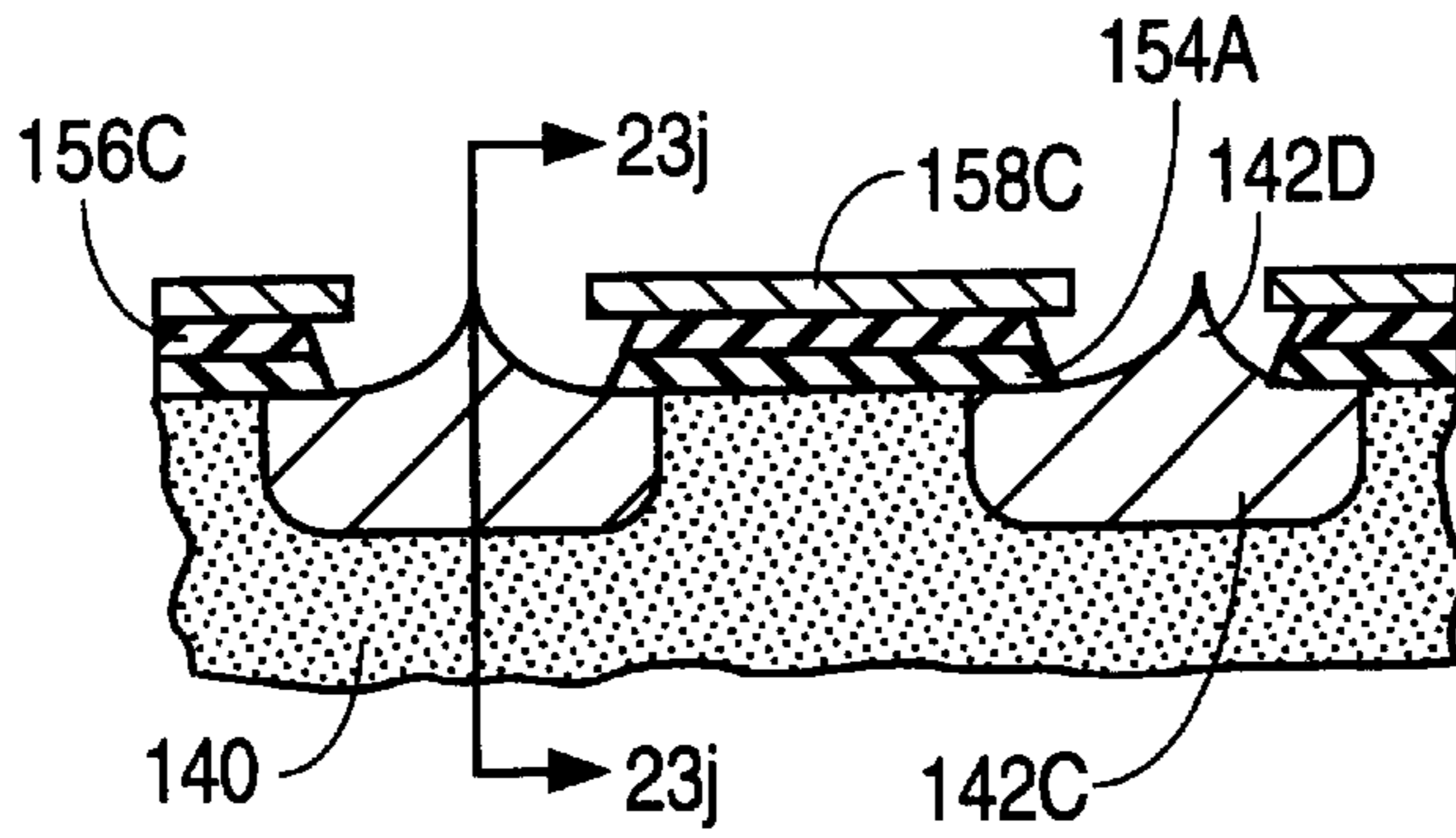


FIG. 25b

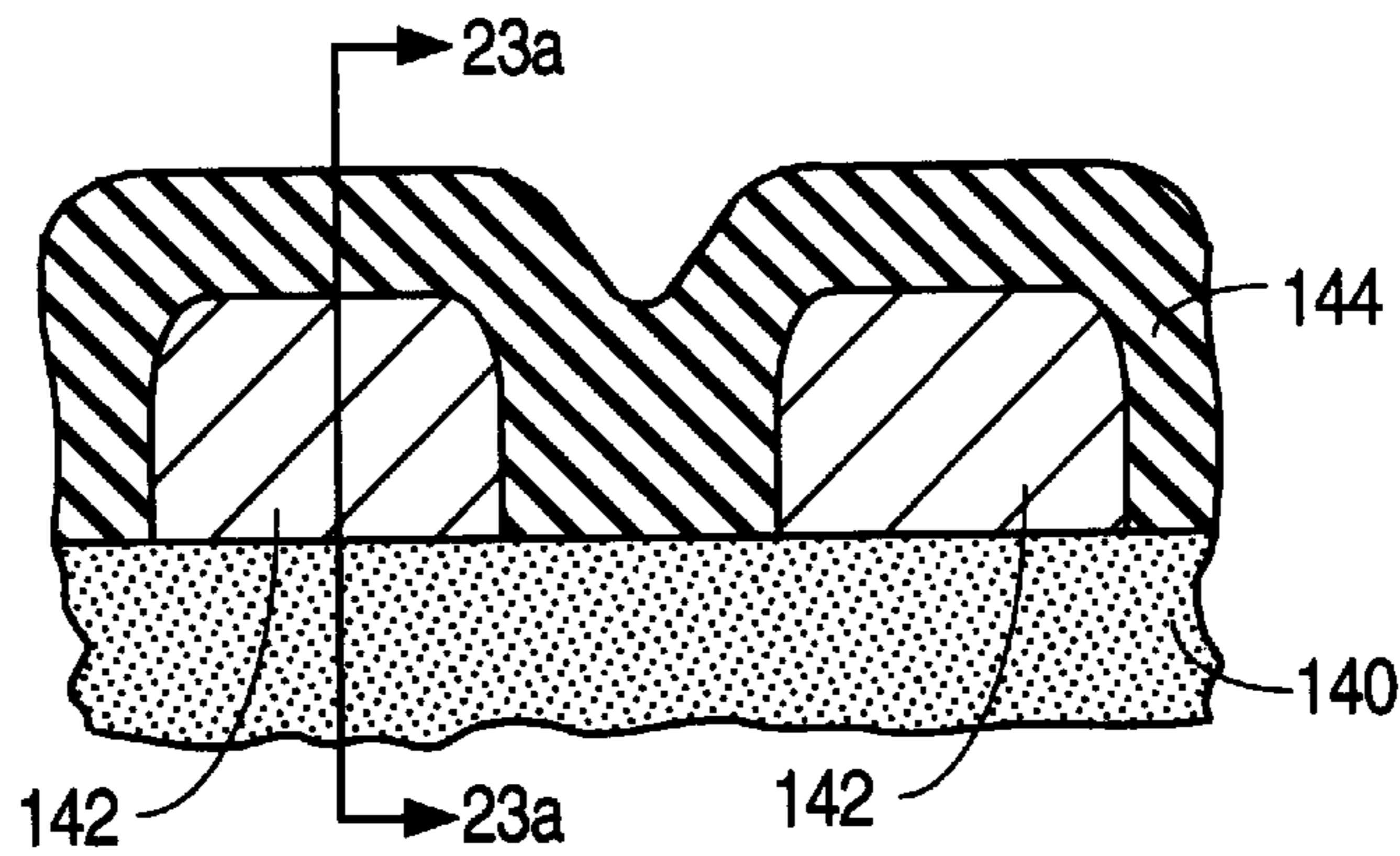


FIG. 26a

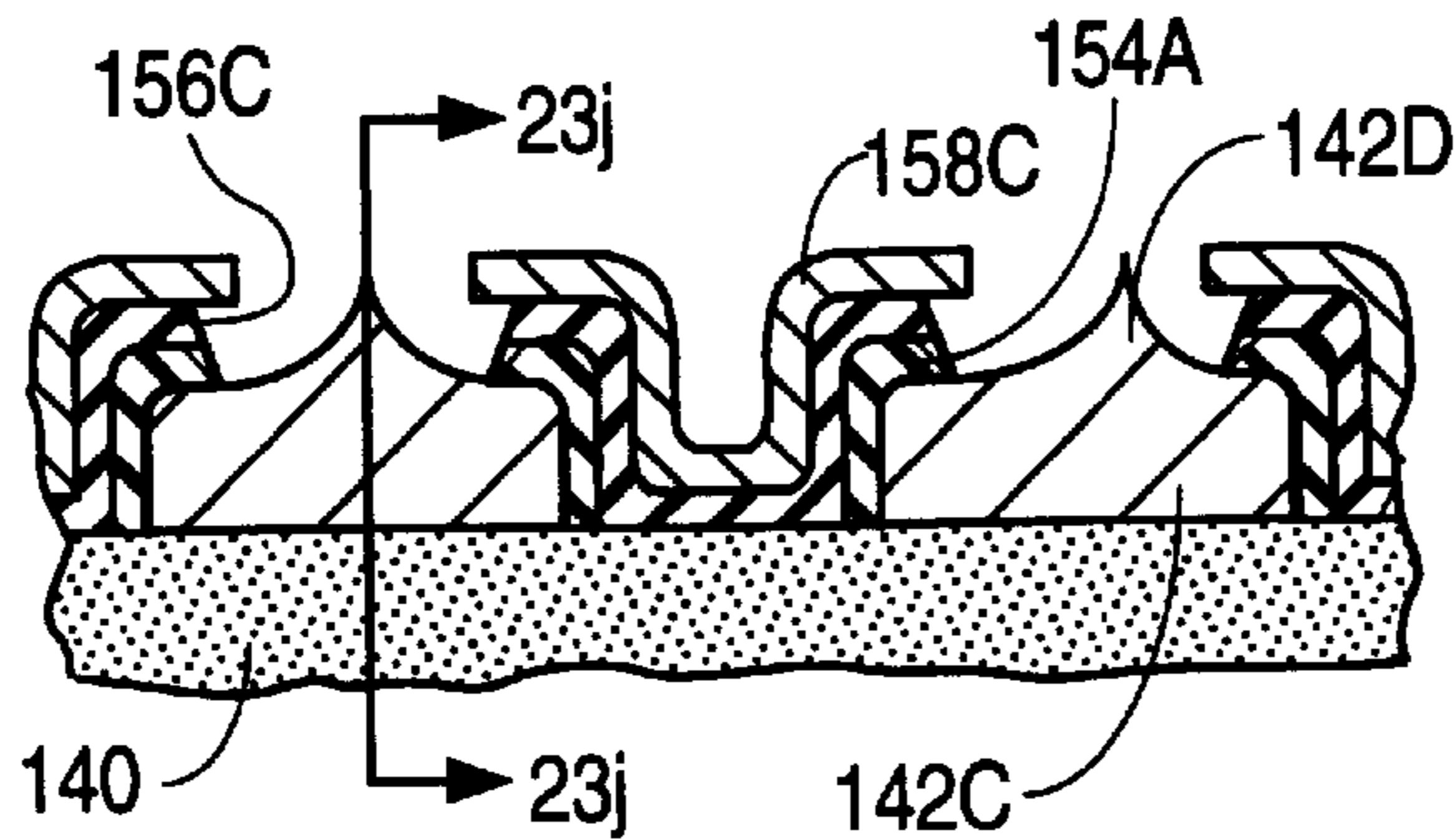


FIG. 26b

USE OF EARLY FORMED LIFT-OFF LAYER IN FABRICATING GATED ELECTRON- EMITTING DEVICES

CROSS-REFERENCE TO RELATED APPLICATIONS

This is a continuation of U.S. patent application Ser. No. 08/269,229, filed 29 Jun. 1994, and now U.S. Pat. No. 5,564,959, which is a continuation-in-part of both U.S. patent application Ser. No. 08/118,490, filed 8 Sep. 1993, now U.S. Pat. No. 5,462,467, and U.S. patent application Ser. No. 08/158,102, filed 24 Nov. 1993, now U.S. Pat. No. 5,559,389, and is related to Spindt et al, co-filed U.S. patent application Ser. No. 08/261,311 "Fabrication of Electron-emitting Structures Using Charged-Particle Tracks and Removal of Emitter Material," filed 29 Jun. 1994. To the extent not repeated herein, the contents of these three related applications are incorporated by reference herein.

FIELD OF USE

This invention relates to electron emission. More particularly, this invention relates to techniques for manufacturing electron-emitting devices, commonly referred to as cathodes, suitable for products such as cathode-ray tube ("CRT") displays of the flat-panel type.

BACKGROUND ART

Cathodes can emit electrons by photoemission, thermionic emission, and field emission, or as the result of negative electron affinity. A field-emission cathode (or field emitter) supplies electrons when subjected to an electric field of sufficient strength. The electric field is created by applying a suitable voltage between the cathode and an electrode, typically referred to as the anode or gate electrode, situated a short distance away from the cathode.

When used in a flat-panel display such as a flat-panel television or video monitor, a field emitter typically contains a group, often a very large group, of individual electron-emissive elements distributed across a supporting structure. This configuration is referred to here as an area field emitter. Busta, "Vacuum microelectronics—1992," *J. Micromech. Microeng.*, Vol. 2, 1992, pp. 43–74, describes a number of different techniques that have been investigated for manufacturing electron-emissive elements in gated area field emitters.

Fischer et al, "Production and use of nuclear tracks: imprinting structure on solids," *Rev. Mod. Phys.*, Oct. 1983, pp. 907–948, describes how nuclear tracks are employed in manufacturing field emitters according to a replica technique. In Fischer et al, nuclear tracks are formed through a substrate. The tracks are etched to create cavities in the substrate after which metal is deposited on the substrate to create a film that extends across the substrate and fills the cavities. The substrate is then removed. The metal film, including the resultant metal protrusions, form an area field emitter as a replica of the substrate.

Some area field emitters employ elongated electron-emissive elements. For example, Yoshida et al, U.S. Pat. No. 5,164,632, discloses a gated field emitter in which solid elongated electron-emissive elements are created in pores extending through a dielectric layer. Greene et al, U.S. Pat. No. 5,150,192, uses hollow elongated electron-emissive elements.

Other area field emitters utilize generally conical electron-emissive elements. See Spindt et al, U.S. Pat. No. 3,665,241.

Also see Borel, U.S. Pat. No. 4,940,916; Betsui, "Fabrication and Characteristics of Si Field Emitter Arrays," *Tech. Dig. IVMC 91*, pp. 26–29; and Fukuta et al, European Patent Publication 508,737 A1.

In yet other area field emitters, electron-emissive particles of various shapes and/or sizes are distributed across a supporting layer at the bottoms of openings that extend through a gate structure overlying the supporting layer. Chason, U.S. Pat. No. 5,019,003, discloses an example of this type of field emitter. Other such examples are disclosed in Thomas et al, U.S. Pat. No. 5,150,019; Jaskie et al, U.S. Pat. No. 5,278,475; and Kane et al, U.S. Pat. No. 5,252,833.

When a portion of a gated area field-emission device in a flat-panel CRT is actively emitting electrons as the result of a suitable applied extraction voltage, the current density produced by emitted electrons ideally should be uniform across the activated portion. In a real field emitter, the emission current density typically becomes more uniform as the emitter packing density—i.e., the number of electron-emission elements per unit area—increases and, correspondingly, as the lateral area occupied by an electron-emissive element decreases.

In manufacturing high-quality prior art gated electron emitters, use of technologies such as photolithography typically places severe restrictions on the minimum lateral size of electron-emissive features such as an electron-emissive element or an opening for an electron-emissive element, especially in a volume production environment. More specifically, depth of field, sometimes referred to as depth of focus, is commonly employed in characterizing radiation-based patterning techniques such as photolithography. Briefly stated, the depth of field is the (maximum) distance, measured along the optic axis, across which an acceptable pattern can be obtained on a generally flat surface situated, generally orthogonal to the optic axis, at any point along that distance.

The depth of field in photolithography is finite and, in particular, is relatively small compared to what would be desirable for efficient manufacturing of area electron emitters on a production scale. Consider an electron-emitting device in which the total area of the surface to be photolithographically patterned is several square centimeters or more. The flatness of the surface being patterned, the presence of features on the surface, and the alignment of the surface in the photolithographic radiation-exposure combined with the small photolithographic depth of field significantly limit the minimum lateral size of features photolithographically defined at the surface using a single radiation exposure.

Finer photolithographic patterns can be obtained by exposing small parts of the total area to the patterning radiation in separate expose-and-move steps. However, such an expose-and-move process is time-consuming and therefore expensive because it requires re-alignment and re-focus before each exposure.

As an example, the conical electron-emissive elements in Betsui and Fukuta et al appear to have a photolithographically defined base diameter of 1–3 μm . It is desirable to overcome these limitations so as to be able to fabricate high-quality gated area electron emitters having smaller lateral electron-emissive features. It is also desirable to increase the emitter packing density so as to attain more uniform emission current density.

GENERAL DISCLOSURE OF THE INVENTION

The present invention furnishes a set of processes for manufacturing gated electron-emitting devices utilizing

charged-particle tracks to define locations for electron-emissive features—e.g., an electron-emissive element or an opening for one or more electron-emissive elements—in the gated electron-emitting device. Use of charged-particle tracks typically enables the lateral areas of the electron-emissive elements to be made quite small. For example, each track-defined electron-emissive element in the invention typically occupies a lateral area having a mean diameter of 0.1–0.2 μm and thus occupies considerably less area than each electron-emissive element in prior art gated electron emitters such as those of Betsui and Fukuta et al.

One item contributing to the small lateral area occupied by an electron-emissive element in the invention is the fact that a charged-particle track constitutes a damaged zone whose mean diameter is typically on the nanometer scale. Furthermore, in contrast to photolithography where the depth of field is finite and, in fact, is relatively small, the depth of field is effectively infinite for charged-particle tracks. As a result, depth of field does not place any significant practical limitations on the minimum lateral emitter feature size attainable in the invention. Creating and etching the charged-particle tracks is no more complex than utilizing photolithography.

An electron emitter fabricated according to the invention is furnished with a gate electrode having gate openings that provide access to the electron-emissive elements. Preferably, the gate openings are self-aligned to the electron-emissive elements or to openings in which the electron-emissive elements are situated. The manufacturing processes of the invention are suitable for commercial volume production of gated area electron-emitting devices for CRT applications such as flat-panel displays. In short, the invention provides a large advance over the prior art.

Specifically, in one aspect of the invention, charged particles are passed through a track layer to provide the track layer with a multiplicity of charged-particle tracks. The track layer is etched along the charged-particle tracks to create corresponding open spaces through the track layer. Electron-emissive elements are then formed at locations respectively centered on the open spaces.

For example, each electron-emissive element can be formed in a corresponding one of the open spaces through the track layer. Formation of the electron-emissive elements thereby broadly entails creating the electron-emissive elements in such a way that they are accessible through the open spaces. Alternatively, the electron-emissive elements can be defined in an emitter region provided below the track layer.

A patterned gate layer is subsequently created over the electron-emissive elements. Gate openings extend through the gate layer in such a way that each gate opening exposes one or more of the electron-emissive elements. Preferably, each gate opening is centered on a corresponding one of the electron-emissive elements. Various processing techniques that key on the electron-emissive elements are employed to create the gate openings in this manner.

In another aspect of the invention, apertures are created through a track layer by forming and etching charged-particle tracks as described above. At this point, the processing sequence diverges from the first-mentioned aspect of the invention in that gate openings are formed through a gate layer underlying the track layer after which electron-emissive elements are created. This is opposite to the first aspect in which the electron-emissive elements are formed before creating gate openings through the gate layer.

For example, the gate openings in the second-mentioned aspect of the invention can be formed at locations centered

on the apertures through the track layer. Material is then deposited through the gate openings in a manner aligned to the gate openings to at least partially form the electron-emissive elements.

A further aspect of the invention likewise begins with creating apertures through a track layer by forming and etching charged-particle tracks in the above-described manner. An underlying electrically non-insulating layer, typically a gate layer, is etched through the apertures to form corresponding further apertures through the non-insulating layer. As discussed further below “electrically non-insulating” means electrically conductive or electrically resistive here.

An underlying electrically insulating layer is then etched through the further apertures, typically gate openings, to form corresponding dielectric open spaces through the insulating layer down to an underlying lower electrically non-insulating region. Electron-emissive elements are typically provided over the lower non-insulating region in such a way that they are electrically coupled to it.

The small size of the electron-emissive elements in an electron emitter fabricated according to the present invention provides a number of advantages. For example, the operating voltages can be considerably lower than that achievable with otherwise equivalent electron emitters of the prior art. Importantly, the emitter packing density can be considerably higher than in the prior art. This produces a much more uniform emission current density across the electron emitter.

The gate layer is typically self-aligned to the electron-emissive features. The size and spacing of the electron-emissive features is not limited by technologies such as photolithography. As a result, the components of the electron emitter can be manufactured at lateral dimensions on the nanometer scale. The fabrication process of the invention is also simple.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1a–1k are cross-sectional front views representing steps in fabricating a gated area field emitter according to the invention.

FIGS. 2a–2k are plan views respectively corresponding to FIGS. 1a–1k. The cross sections of FIGS. 1a–1k are taken through stepped plane 1—1 in FIGS. 2a–2k.

FIGS. 3a–3f are cross-sectional front views representing an alternative set of steps performable on the structure of FIGS. 1f and 2f in fabricating a gated area field emitter according to the invention.

FIGS. 4a–4f are plan views respectively corresponding to FIGS. 3a–3f. The cross sections of FIGS. 3a–3f are taken through stepped plane 3—3 in FIGS. 4a–4f.

FIGS. 5a–5j are cross-sectional front views representing steps in fabricating a gated area field emitter according to the invention.

FIGS. 6a–6j are plan views respectively corresponding to FIGS. 5a–5j. The cross sections of FIGS. 5a–5j are taken through stepped plane 5—5 in FIGS. 6a–6j.

FIGS. 7a–7e are cross-sectional front views representing an alternative set of steps performable on the structure of FIGS. 5i and 6i for manufacturing a gated area field emitter according to the invention.

FIGS. 8a and 8b are respective expanded cross-sectional front views of one of the electron-emissive elements in FIGS. 7c and 7e.

FIGS. 9a–9h are cross sectional front views representing an alternative set of steps performable on the structure of

FIGS. 5f and 6f for manufacturing a gated area field emitter according to the invention.

FIGS. 10a–10e are cross-sectional front views representing a further set of steps performable on the structure of FIGS. 5f and 6f for manufacturing a gated area field emitter according to the invention.

FIGS. 11a–11e are plan views respectively corresponding to FIGS. 10a–10e. The cross sections of FIGS. 10a–10e are taken through plane 10—10 in FIGS. 11a–11e.

FIGS. 12a–12j are cross-sectional front views representing steps in fabricating a gated field emitter according to the invention.

FIGS. 13.1 and 13.2 are cross-sectional front and side views depicting how the initial structure of FIG. 1a or 5a appears when the lower non-insulating region consists of an electrically conductive part and an electrically resistive part. The cross section of FIG. 13.1 is taken through plane 13.1—13.1 in FIG. 13.2. The cross section of FIG. 13.2 is taken through plane 13—13 in FIG. 13.1.

FIGS. 14.1–14.4 are cross-sectional front views respectively depicting how the final structures of FIGS. 5j, 7e, 9h, and 10e appear when they contain the two-part lower non-insulating region of FIG. 12.

FIGS. 15.1 and 15.2 are expanded cross-sectional front views of alternative electron-emissive elements employable in the present field emitters where each electron-emissive element consists of an electrically resistive portion and an overlying electron-emissive portion.

FIGS. 16.1 and 16.2 are expanded cross-sectional front views of alternative shapes for electron-emissive cones in the present field emitters.

FIGS. 17.1–17.4 are cross-sectional longitudinal views of differently shaped electron-emissive filaments usable in the field-emission structures of FIGS. 1k, 3f, and 5j.

FIG. 18 is a functional diagram for the components of an electrochemical deposition system used in the invention.

FIGS. 19a–19d are cross-sectional front views representing another set of steps performable on the structure of FIGS. 5f and 6f for manufacturing a gated area field emitter according to the invention.

FIGS. 20a–20d are plan views respectively corresponding to FIGS. 19a–19d. The cross sections of FIGS. 19a–19d are taken through stepped plane 19—19 in FIGS. 20a–20d.

FIGS. 21a–21e are cross-sectional front views representing yet another set of steps performable on the structure of FIGS. 5f and 6f for manufacturing a gated field emitter according to the invention.

FIGS. 22a–22c are cross-sectional front views representing a set of steps performable on a variation of the structure in FIGS. 5f and 6f for manufacturing a gated area field emitter according to the invention.

FIGS. 23a–23j, are cross-sectional front views representing steps in fabricating a gated area field emitter according to the invention.

FIGS. 24a–24d are plan views respectively corresponding to FIGS. 23b, 23c, 23f, and 23j. The cross sections of FIGS. 23b, 23c, 23f, and 23j are respectively taken through planes 23b—23b, 23c—23c, 23f—23f, and 23j—23j in FIGS. 24a–24d.

FIGS. 25a and 25b are cross-sectional side views respectively corresponding to FIGS. 23a and 23j for an embodiment in which the emitter lines are conductively doped regions formed in an electrically resistive semiconductor substrate. The cross sections of FIGS. 23a and 23j are

respectively taken through planes 23a—23a and 23j—23j in FIGS. 25a and 25b. The cross section of FIG. 25a is taken through plane a—a in FIG. 23a. The cross section of FIG. 25b is taken through stepped plane b—b in FIGS. 23j and 24d.

FIGS. 26a and 26b are cross-sectional side structural views respectively corresponding to FIGS. 23a and 23j for an embodiment in which the emitter lines consist of metal or conductively doped semiconductor material formed on an electrically insulating or resistive substrate. The cross sections of FIGS. 23a and 23j are respectively taken through planes 23a—23a and 23j—23j in FIGS. 26a and 26b. The cross section of FIG. 26a is taken through plane a—a in FIG. 23a. The cross section of FIG. 26b is taken through stepped plane b—b in FIGS. 23j and 24d.

Like reference symbols are employed in the drawings and in the description of the preferred embodiments to represent the same, or very similar, item or items.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The following definitions are used in the description below. The “mean diameter” for a two-dimensional item of non-circular shape is the diameter of a circle of the same area as the non-circular item. The “mean diameter” for a three-dimensional item of non-spherical shape either is the diameter of a sphere of the same volume as the non-spherical item or is the diameter of a right circular cylinder of the same volume as the item. The equal-volume cylinder diameter is generally used when the item is cylindrical or considerably elongated.

Herein, the term “electrically insulating” (or “dielectric”) generally applies to materials having a resistivity greater than 10^{10} ohm-cm. The term “electrically non-insulating” thus refers to materials having a resistivity below 10^{10} ohm-cm. Electrically non-insulating materials are divided into (a) electrically conductive materials for which the resistivity is less than 1 ohm-cm and (b) electrically resistive materials for which the resistivity is in the range of 1 ohm-cm to 10^{10} ohm-cm. These categories are determined at an electric field of no more than 1 volt/ μ m.

Examples of electrically conductive materials (or electrical conductors) are metals, metal-semiconductor compounds (such as metal silicides), and metal-semiconductor eutectics (such as gold-germanium). Electrically conductive materials also include semiconductors doped (n-type or p-type) to a moderate or high level. Electrically resistive materials include intrinsic and lightly doped (n-type or p-type) semiconductors. Further examples of electrically resistive materials are cermet (ceramic with embedded metal particles), other such metal-insulator composites, graphite, amorphous carbon, and modified (e.g., doped or laser-modified) diamond.

Referring to the drawings, FIGS. 1a–1k (collectively “FIG. 1”) and FIGS. 2a–2k (collectively “FIG. 2”) illustrate a process for manufacturing a gated field-emission cathode structure using charged-particle tracks according to the teachings of the invention. The field-emission structure is typically used to excite phosphors on a face plate in a CRT of a flat-panel display such as a flat-panel television or a flat-panel video monitor for a personal computer, a lap-top computer, or a work station.

The starting point for the fabrication process is an electrically insulating substrate 20 typically consisting of ceramic or glass. Substrate 20 is typically configured as a plate having a largely flat upper surface and a largely flat

lower surface substantially parallel to the upper surface. In a flat-panel CRT display, substrate **20** constitutes at least part of the backplate (or baseplate).

Substrate **20** furnishes support for the field-emission structure. As such, the substrate thickness is at least $500\ \mu\text{m}$. In a 25-cm (diagonal) flat-panel CRT display where internal supports are placed between the phosphor-coated face plate and the field emitter, the substrate thickness is typically 1–2 mm. If substrate **20** provides substantially the sole support for the field emitter, the substrate thickness is typically 4–14 mm.

A lower electrically non-insulating region **22** is formed along the top of substrate **20** as indicated in FIGS. **1a** and **2a**. Lower non-insulating region **22** usually consists of an electrical conductor, preferably a metal such as chromium. In this case, the thickness of region **22** is $0.05\text{--}1.5\ \mu\text{m}$. Other candidates for region **22** include tantalum, tungsten, nickel, and molybdenum. Region **22** could also be formed with (a) conductively doped (i.e., moderately or heavily doped) semiconductor material such as n-type or p-type silicon, (b) metal-semiconductor compounds such as metal silicides, or/and (c) metal-semiconductor eutectics such as gold-germanium.

Lower non-insulating region **22** is typically a patterned layer containing a group of parallel lines, of which two such lines are depicted in FIG. **2a**. When region **22** is configured in this way, the final field-emission structure is particularly suitable for selectively exciting phosphors in a flat-panel display. These lines are typically no more than $300\ \mu\text{m}$ wide for a 25-cm flat-panel display. Nonetheless, region **22** can be arranged in various other patterns, or can even be unpatterned.

A largely homogeneous electrically insulating track (or track-recording) layer **24** is formed on the top of the structure. See FIGS. **1b** and **2b**. Parts of insulating track layer **24** are situated on both substrate **20** and lower non-insulating region **22**. The thickness of layer **24** is $0.1\text{--}2\ \mu\text{m}$. Suitable dielectric materials for track layer **24** fall into three groups: (a) organic polymers such as polycarbonate, polystyrene, and cellulose acetate, (b) inorganic glasses such as phosphate, silicate, soda-lime, and spin-on glasses, and (c) crystals such as mica and quartz. Preferably, layer **24** consists of inorganic glass having a thickness of $1\ \mu\text{m}$.

Insulating track layer **24** is subjected to energetic charged particles that impinge on the top of layer **24** in a direction largely perpendicular to the lower surface of substrate **20** and thus in a direction generally perpendicular to the upper structural surface. The charged particles have sufficient energy to form straight tracks through layer **24** at random locations across layer **24**. The charged-particle tracks constitute damaged zones created along the paths of the charged particles. Each track has a heavily damaged core whose mean diameter is in the vicinity of 4 nm. As shown in FIGS. **1c** and **2c**, the tracks consist of (a) charged-particle tracks **26₁** through the portions of layer **24** overlying non-insulating region **22** and (b) charged-particle **26₂** through the portions of layer **24** directly overlying substrate **20**.

Charged-particle tracks **26₁** and **26₂** (collectively “**26**”) extend parallel to one another in a direction generally perpendicular to the upper structural surface. Although charged-particle tracks **26** are randomly distributed across insulating track layer **24**, they have a well-defined average spacing. The track density is usually in the range of $10^6\text{--}10^9$ tracks/cm². A typical value falls in the narrower range of $10^7\text{--}10^8$ tracks/cm² which yields an average track spacing of approximately 1–3 μm . For illustrative purposes, only a very small portion of tracks **26** are indicated in FIGS. **1c** and **2c**.

The charged particles penetrate into the material below insulator **24**. Such penetration is not material here and, accordingly, is not shown in the drawings.

In a typical implementation, a charged-particle accelerator which forms a well-collimated beam of ions is employed to form tracks **26**. The ion beam is scanned uniformly across insulating track layer **24**. The preferred charged-particle species is doubly ionized argon (Ar^{++}) at an energy of 8 MeV. Alternatively, tracks **26** could be created from a collimated source of nuclear fission particles produced, for example, by the radioactive element californium **252**.

Insulating track layer **24** is brought into contact with (e.g., by immersion) a suitable chemical etchant that attacks the damaged insulating material along tracks **26** much more than the undamaged material of layer **24**. Pores **28₁** and **28₂** are thereby etched through layer **24** respectively along tracks **26₁** and **26₂**. See FIGS. **1d** and **2d**. For illustrative purposes, the lateral dimensions of pores **28₁** and **28₂** (collectively “**28**”) compared to the widths of the lines that form conductive layer **20** are greatly exaggerated in the plan-view drawings. The remainder of layer **24** now constitutes homogeneous porous insulating track layer **24A**.

Pores **28** are generally circular in shape as viewed from the top in the plan view of FIG. **2d**. Depending on how the track etching is done, pores **28** can be cylindrical or (slightly) conical in three dimensions. For purposes of illustration, pores **28** are represented as cylinders in the drawings. The pore diameter can vary from 4 nm to $2\ \mu\text{m}$. Preferably, the pore diameter lies in the range of 10–200 nm. A typical value is 100 nm.

The etchant used for creating pores **28** preferably does not significantly attack substrate **20** or non-insulating region **22**. When components **20**, **22**, and **24A** respectively consist of ceramic, chromium, and polycarbonate, the etching is done in 6.25 normal sodium hydroxide at 50°C . The etch time is less than 10 minutes for the typical $1\text{-}\mu\text{m}$ track-layer thickness.

Solid electron-emissive metal filaments **30** are formed in pores **28₁**, which overlie non-insulating region **22**, by electrochemically depositing a suitable filament material. See FIGS. **1e** and **2e**. The deposition is performed in an electrochemical cell, described below, in which non-insulating region **22** acts as a deposition cathode. After bringing the structure into contact with the cell electrolyte and activating the cell, current flows between cathode **22** and a separate anode. The filament material uniformly builds up in pores **28₁** starting from non-insulating region **22**. The lower ends of filaments **30** contact region **22**.

Substantially none of the filament material accumulates in pores **28₂** directly above substrate **20** because there is no electrical contact at the bottoms of pores **28₂**. In FIG. **2e**, the dark circles represent filaments **30** in filled pores **28₁**, while the light circles represent empty pores **28₂**. The use of conductive layer **22** as the cathode during the electrochemical deposition enables the deposition to be selective.

The electrochemical deposition is typically performed for a time sufficient to enable the upper ends of pores **30** to be nearly coplanar with the top of porous track layer **24A**. This situation is illustrated in FIG. **1e**. The deposition can, however, be performed for a longer time so that filaments **30** bulge slightly out of pores **28₁** or for a shorter time so that the upper ends of filaments **30** are significantly below the top of layer **24A**.

The upper ends of filaments **30** are preferably formed with a noble metal. In a typical implementation, the upper ends of filaments **30** consist of platinum when non-insulating region

22 is chromium. Other candidate noble metals for the upper filament ends are gold and palladium. The remaining portions of filaments **30** typically consist of the same (noble) metal as the upper ends but can be formed with another metal such as nickel, copper, cobalt, molybdenum, or niobium.

Elements **30** are true filaments for which the ratio of length to maximum diameter is at least 2 and normally at least 3. The length-to-maximum-diameter ratio is preferably 5 or more. Filaments **30** are typically cylinders of circular transverse cross section. Nonetheless, the transverse cross section can be somewhat non-circular. In any case, the ratio of maximum diameter to minimum diameter for each filament **30** is usually no more than 2.

Filaments **30** are all of substantially the same length. The filament length is 0.1–2 μm , typically 1 μm . In this regard, the average track spacing and, consequently, the average filament spacing are arranged to be somewhat greater than the filament length.

In order to create a gate electrode for the structure, electrically conductive caps **32** are electrochemically deposited respectively on filaments **30**. See FIGS. 1*f* and 2*f*. Caps **32** are generally circular in shape as viewed from the top in FIG. 2*f*.

Each cap **32** is centered on the upper end of corresponding filament **30**. Along the upper surface of track layer **24A**, each cap **32** also reaches a greater diameter than underlying filament **30**. As a result, each cap **32** has a lateral periphery that encloses the lateral periphery of underlying filament **30** along the top of layer **24A**. Typically, the average diameter of caps **30** along the top of layer **24A** is at least 1.5 times the average diameter of filaments **30**.

The electrochemical deposition to create caps **32** is performed in an electrochemical cell, again described below, in which filaments **30** (attached to non-insulating region **22**) act as a cathode. After the structure has been brought into contact with the cell electrolyte and the cell has been activated, current flows between filaments **30** and a separate anode. The cap material thereby builds up on filaments **30** until the desired cap diameter is reached.

Caps **32** consist of a metal different from the filament metal along the upper ends of filament **30**. In particular, the cap metal is selectively etchable with respect to the directly underlying filament metal. When filaments **30** consist of platinum, caps **32** are formed with a metal such as silver whose half-cell potential is less positive than that of platinum.

Next, part of the thickness of porous track layer **24A** is uniformly removed along the top of layer **24A** to produce the structure shown in FIGS. 1*g* and 2*g*. Item **24C** is the remainder of porous track layer **24A**. As illustrated in FIG. 1*g*, caps **32** are vertically separated from remaining track layer **24C**.

The track-material removing step is performed with an etchant that does not significantly attack the filament or cap metal. Likewise, the etchant does not significantly attack substrate **20** or non-insulating region **22**. The etching is conducted for a time sufficient to remove approximately 0.3 μm of the thickness of the track material.

Electrically non-insulating gate material is now deposited on top of the structure to a thickness less than the removed thickness of the porous material. The gate-material thickness is typically 0.1 μm . The deposition is performed in a direction largely perpendicular to the lower surface of substrate **20** and thus in a direction generally perpendicular to the upper structural surface. As indicated in FIGS. 1*h* and

2*h*, a layer **34A** of the gate material thereby accumulates on the portion of track layer **24C** not shielded by caps **32**. A layer **34B** of the gate material likewise normally accumulates on each cap **32**. Importantly, caps **32** prevent substantially any of the gate material from accumulating on the portions of track layer **24C** below caps **32**.

The criteria for selecting the gate material normally an electrical conductor, depends on the technique utilized below for removing caps **32**. If the cap removal is done electrochemically, non-insulating layer **34A** which later becomes the gate electrode can be electrically protected. As a result, the gate material can generally be any metal that is not highly reactive. Suitable candidates include molybdenum, copper, and aluminum.

If chemical or plasma etching is used for the cap removal, the gate material consists of a metal different from the cap metal. In particular, the cap metal must be selectively etchable with respect to the gate material. The gate material could also be conductively doped semiconductor material, provided that the selective etchability requirement is, as appropriate, met.

An evaporative deposition technique is typically used to form non-insulating layers **34A** and **34B**. The evaporative deposition is performed at low pressure in a suitable vacuum chamber. Layers **34A** and **34B** could also be formed by other direction-controllable physical deposition techniques such as sputtering.

Caps **32** are removed with an etchant that attacks the cap metal much more than the gate material or the filament metal at the upper ends of filaments **30**. In so doing, non-insulating portions **34B** are simultaneously removed. When regions **30**, **32**, and **34A** respectively consist of platinum, silver, and molybdenum, the etching is typically performed by an electrochemical process in which non-insulating layer **34A** is held at a different potential than caps **32**. This is achieved by applying one potential to non-insulating layer **34A** and another potential to lower non-insulating region **22**. Alternatively, as mentioned above, caps **32** and overlying non-insulating portions **34B** can be removed with a chemical or plasma etchant.

FIGS. 1*i* and 2*i* show the resultant structure in which the upper ends of filaments **30** are now exposed. Patterned upper non-insulating layer **34A** now constitutes the gate electrode for the field-emission structure. Gate electrode **34A** has gate openings **36** respectively centered on filaments **30**. Due to the protection supplied by caps **32** during the gate-material deposition, each gate opening **36** is wider than corresponding filament **30**. Consequently, gate electrode **34A** is laterally separated from filaments **30**.

The cathode/gate structure of FIGS. 1*i* and 2*i* can be utilized directly as a field emitter. Nonetheless, it is advantageous for filaments **30** to extend out of the track material and for the upper ends of filaments **30** to be sharpened.

To this end, portions of porous track layer **24C** exposed through openings **36** are removed with an isotropic etchant to form cavities **38** respectively around filaments **30**. See FIGS. 1*j* and 2*j*. The isotropic etchant used to create cavities **38** does not significantly attack any other portions of the structure. Filaments **30** now protrude outward beyond the remainder **24D** of porous track **24C**. Each cavity **38** is considerably wider—i.e., has a considerably greater maximum transverse cross-sectional area—than corresponding pore **28₁**.

Cavities **38** typically extend partway down to lower non-insulating region **22**. This situation is illustrated in FIG. 1*j*. However, cavities **38** can extend all the way down to

region 22. In either case, by appropriately choosing the fabrication parameters, track layer 24D is sufficient for supporting gate layer 34A.

An electropolishing and etching operation is performed to tailor and sharpen the upper ends of filaments 30. The electropolishing, which is done first, rounds the upper ends of filaments 30 and reduces their length somewhat. The etch sharpens the rounded filament ends.

The electropolishing is performed in an electrolytic cell in which filaments 30 constitute the anode. After bringing the structure of FIGS. 1j and 2j into contact with the cell electrolyte, a suitable potential is applied between (a) filaments 30 by way of non-insulating region 22 and (b) a cathode plate, typically gate layer 34A, to cause current to flow between filaments 30 and the cathode plate. The electropolishing is typically conducted in an operational regime where the rate of metal removal increases with increasing electric field strength. Because the highest electric field, and thus the greatest material removal rate, for each filament 30 occurs at the outer top filament edge, the upper ends of filaments 30 become rounded.

By arranging the lower surface of substrate 20 to be approximately parallel to the cathode plate in the electrolytic cell, the upper ends of the longer ones of filaments 30 experience the highest electric fields. More material is thereby removed from the upper ends of the longer ones of filaments 30 than from the shorter ones during the electropolishing. Accordingly, the electropolishing also enables filaments 30 to become more uniform in length.

The filament etch is also typically done by bringing the field-emission structure into contact with the electrolyte of an electrolytic cell in which filaments 30 constitute the anode to which a driving voltage is applied via non-insulating region 22. The rate of metal removal varies with electric field strength and filament morphology in such a way that the rounded upper ends of filaments 30 become pointed. The etch can also be done according to a chemical technique. If the etch is done by a chemical technique rather than an electrochemical technique, undesired etching of gate layer 34A can be avoided by applying an appropriate voltage to layer 34A and/or making suitable materials selection. FIGS. 1k and 2k illustrate the final field emitter in which sharpened filaments 30B are the remainders of filaments 30.

An electropolishing step is also typically performed to round the edges of patterned gate layer 34A overlying cavities 38. Layer 34A acts as the anode during this step, while filaments 30B serve as the cathode via non-insulating region 22. Patterned gate layer 34B in FIGS. 1k and 2k is the edge-rounded remainder of gate layer 34A.

FIGS. 3a-3f (collectively "FIG. 3") and FIGS. 4a-4f (collectively "FIG. 4"), depict an alternative group of steps for providing the cathode structure of FIGS. 1f and 2f with a self-aligned gate structure. FIGS. 1f and 2f are repeated here as FIGS. 3a and 4a. In this alternative, the thickness of porous track layer 24A is typically 0.5 μm .

A blanket layer 40 of electrically non-insulating gate material is deposited on porous track layer 24A and caps 32 along the top of the structure. See FIGS. 3b and 4b. The thickness of layer 40 is typically 0.1 μm .

The gate material usually consists of an electrical conductor, preferably a metal selectively etchable with respect to the cap metal. Platinum can be utilized for the gate metal. As long as the selective etchability requirement is met, the gate material could consist of other metals or/and conductively doped semiconductor material. Various techniques can be employed for the gate-material deposition

provided that the thickness of the portion of gate-material layer 40 above track layer 24A is relatively uniform.

The portions of gate-material layer 40 overlying caps 32 are removed in a planarization operation by which the combination of caps 32 and layer 40 is provided with a largely flat upper surface. In so doing, small upper portions of caps 32 are also removed. FIGS. 3c and 4c depict the resulting structure in which items 32A are the remainders of caps 32. Item 40A is the remainder of gate-material layer 40.

The planarization operation is performed in several steps. Firstly, a flowable material such as photoresist is deposited on the top of the structure. Secondly, the flowable material is flowed so that its upper surface becomes largely planar. Thirdly, an etchback is performed with an etchant that attacks the flowable material and the gate material at approximately the same rate. The etchback is terminated when all of the flowable material has been removed. Because the etchant attacks the gate material at approximately the same rate as the flowable material, the upward-protruding portions of gate-material layer 40—i.e., the portions overlying caps 32—are simultaneously removed.

Caps 32A are removed with an etchant that does not significantly attack filaments 30 or non-insulating layer 40A. See FIGS. 3d and 4d. Layer 40A, which now constitutes the gate electrode for the structure, has gate openings 42 respectively centered on filaments 30. Because each cap 32 was wider than corresponding filament 30, gate layer 40A is laterally separated from filaments 30.

As with the cathode/gate structure of FIGS. 1i and 2i, the cathode/gate structure of FIGS. 3d and 4d can be employed directly as a field emitter. However, it is similarly advantageous to further process the structure of FIGS. 3d and 4d in the same manner as that of FIG. 1i and 2i. Accordingly, portions of porous track layer 24A exposed through openings 42 are removed with an isotropic etchant to form cavities 44 around filaments 30 as shown in FIGS. 3e and 4e. Filaments 30 protrude outward beyond the remainder 24E of porous track layer 24A. Each cavity 44 is considerably wider than corresponding pore 28₁.

Cavities 44 typically extend partway down to non-insulating region 22. FIG. 3e illustrates this situation. As in the fabrication process of FIGS. 1 and 2, cavities 44 can also extend fully down to region 22.

An electropolishing and etching operation is performed to adjust and sharpen the upper ends of filaments 30. See FIGS. 3f and 4f. Items 30B again are the sharpened remainders of filaments 30. An additional electropolishing step is typically also done to round the edges of patterned gate layer 40A overlying cavities 44. Item 40B in FIGS. 3f and 4f is the edge-rounded remainder of gate layer 40A.

FIGS. 5a-5j (collectively "FIG. 5") and FIGS. 6a-6j (collectively "FIG. 6") illustrate a further process for manufacturing a gated field-emission structure using charged-particle tracks according to the invention. This field emitter is structurally similar to both that of FIGS. 1k and 2k and that of FIGS. 3f and 4f. Likewise, the field emitter fabricated according to the process of FIGS. 5 and 6 is typically used for exciting phosphors in a flat-panel CRT display.

The starting point again is insulating substrate 20 over which non-insulating region 22 is provided. See FIGS. 5a and 6a. Substrate 20 and region 22 have the characteristics given above. In particular, region 22 typically contains a group of largely parallel lines, two of which are shown in FIG. 6a.

Insulating track layer 24 is again formed on top of the structure. See FIGS. 5b and 6b. Layer 24 likewise has the

characteristics given above since charged-particle tracks are later formed through it.

At this point, the process of FIGS. 5 and 6 deviates from the earlier-described fabrication processes of the invention. A blanket electrically non-insulating layer 46, which later becomes the gate electrode, is formed on the top of insulating layer 24 as indicated in FIG. 5b. Non-insulating layer 46 is typically 0.05 μm in thickness. Layer 46 consists of a metal such as molybdenum, copper, or aluminum. Layer 46 could also be formed with conductively doped semiconductor material. The method for creating layer 46 is not particularly critical as long as its thickness is relatively uniform. Layer 46 is typically formed by a physical vapor deposition technique.

A further electrically insulating track (or track-recording) layer 48 is formed on the top of non-insulating layer 46. Again see FIG. 5b. Insulating track layer 48 is in the range of 0.2–0.5 μm in thickness. As with insulating layer 24, layer 48 consists of (a) an organic polymer such as polycarbonate, polystyrene, or cellulose acetate, (b) an inorganic glass such as phosphate, silicate, soda-lime, or spin-on glass, or (c) a crystal such as mica or quartz. Although there are some implementations in which layer 48 consists of the same insulator as layer 24, layer 48 is normally formed with an insulator that is selectively etchable with respect to layer 24.

The structure is now subjected to energetic charged particles that impinge on the top of insulating track layer 48 in a direction largely perpendicular to the (unshown) flat lower surface of substrate 20 and thus in a direction generally perpendicular to the upper structural surface. The charged particles pass through layers 48, 46, and 24 and into the underlying material to form straight tracks through track layers 24 and 48 at random locations. FIGS. 5c and 6c illustrate the track formation. The charged-particle tracks again constitute damaged zones along the particle paths.

The charged-particle tracks are indicated by reference symbols beginning with “50” in FIGS. 5c and 6c. Each track is divided into (a) a “50A” segment extending through insulating layer 24 and (b) a “50B” segment extending through insulating layer 48 in a straight line with the 50A segment. Although the charged particles pass through non-insulating layer 46, they do not significantly damage layer 46 and therefore do not create charged-particle tracks through layer 46. The tracks fall into two categories: (a) segments 50A₁ and 50B₁ (collectively “50₁”) extending respectively through portions of layers 24 and 48 overlying non-insulating region 22 and (b) segments 50A₂ and 50B₂ (collectively “50₂”) extending respectively through portions of layers 24 and 48 not overlying region 22.

As with charged-particle tracks 26 in the previous fabrication processes of the invention, charged-particle tracks 50₁ and 50₂ (collectively “50”) extend parallel to one another in a direction generally perpendicular to the upper structural surface. Tracks 50 have the same characteristics—e.g., size and spacing—as tracks 26. Tracks 50 are also formed in the same way as tracks 26. To simplify the illustration, only a small portion of tracks 50 are indicated in FIGS. 5c and 6c.

The damaged insulating material along track segments 50B in insulating track layer 48 is removed by bringing layer 48 into contact with a suitable chemical etchant that attacks the damaged 50B track material much more than the undamaged material of layer 48. As a result, generally circular pores are etched through layer 48 along segments 50B down to non-insulating layer 46. The etchant preferably does not significantly attack any of the other parts of the field-emission structure.

The etch is continued into the largely undamaged material of insulating track layer 48 to broaden the pores. Apertures 52₁ and 52₂ are thereby respectively created along track segments 50B₁ and 50B₂. See FIGS. 5d and 6d. Apertures 52₁ and 52₂ (collectively “52”) expose corresponding portions of the upper surface of non-insulating layer 46. The etch is performed in a laterally uniform manner. Accordingly, each aperture 52 is centered on the location of corresponding track segment 50B. The thickness of layer 48 is also reduced during the etch.

The second part of the insulating-material etch can be done with the etchant used during the first part or with another etchant. In either case, components 20, 22, 24, and 46 are not significantly attacked during the second part of the etch. Apertures 52 thereby reach an average mean diameter of 15–300 nm, typically 140 nm, along the bottom of the reduced-thickness remainder 48A of insulating track layer 48. The aperture diameter is substantially the same for all of apertures 52.

Apertures 52 are generally circular in shape as viewed from the top in the plan view of FIG. 6d. Apertures 52 are also usually somewhat wider at the top than at the bottom as indicated in FIG. 5d. The reason for slanting apertures 52 in this way is to facilitate subsequent reactive-ion etching of non-insulating layer 46. The aperture slope is obtained by using etchant whose selectivity—i.e., ratio of the rate at which the damaged track material is attacked to the rate at which the undamaged track material is attacked—is relatively low—e.g., 10 or less.

Apertures 52₂, which are formed as a by-product of creating apertures 52₁, do not serve any useful purpose in the emitter fabrication process. If desired, the formation of apertures 52₂ and the additional (non-useful) features subsequently produced as a result of apertures 52₂ can be suppressed. For example, during the etching of track layer 48, a suitable mask can be utilized to cover track segments 50B₂.

Using track layer 48A as an etch mask, the portions of non-insulating layer 46 exposed through apertures 52 are removed with an anisotropic etchant to create generally circular openings down to track layer 24. FIGS. 5e and 6e depict the consequent structure in which gate openings 54₁ extend through the portions of layer 46 located above non-insulating region 22, while further gate openings 54₂ extend through the portions of layer 46 not overlying region 22. The remainder 46A of layer 46 is the patterned gate electrode for the field emitter. The anisotropic etch is typically done according to a reactive-ion-etch technique.

The etchant utilized to create openings 54₁ and 54₂ (collectively “54”) is controlled in such a way as to avoid significantly attacking other parts of the structure. Due to the anisotropic nature of the etch, each opening 54 is of largely the same transverse shape—i.e. generally circular—and of nearly the same diameter as corresponding aperture 52 (along the bottom of track layer 48a). Each opening 54 is vertically aligned with corresponding aperture 52. Since each aperture 52 is centered on the location of corresponding track segment 50B, each opening 54 is also centered on the location of corresponding segment 50B.

Track layer 48A is usually removed at some point subsequent to the creation of openings 54. As, for example, indicated in FIGS. 5f and 6f, layer 48A could be removed directly after openings 54 are formed.

With portions of the upper surface of insulating track layer 24 now exposed at track segments 50A, the structure is brought into contact with a chemical etchant that attacks

the damaged material along segments **50A** much more than the undamaged material of layer **24**. Pores **28₁** and **28₂** are thereby created through layer **24** respectively along track segments **50A₁** and **50A₂** as indicated in FIGS. **5g** and **6g**. Pores **28₁** and **28₂** (again collectively “**28**”) have the same physical/spatial characteristics, described above, as pores **28** in the previous manufacturing processes of the invention. The etch utilized to create pores **28** here is typically performed in the same manner as described above in connection with the structure of FIGS. **1d** and **2d**. The remainder of layer **24** again is homogeneous porous insulating track layer **24A**.

Importantly, each pore **28** in FIGS. **5g** and **6g** is considerably narrower than corresponding opening **54**. For example, when the bottom diameter of one of apertures **52** is 140 nm so that corresponding opening **54** is 150 nm in diameter, corresponding pore **28** typically has a diameter of 50–100 nm. Each pore **28** is centered on the location of corresponding track segment **50A**. Because (a) each track segment **50B** was in a straight line with corresponding track segment **50A** and (b) each opening **54** is centered on the location of corresponding track segment **50B**, each opening **54** is centered on corresponding pore **28**.

A suitable filament metal is now electrochemically deposited to form electron-emissive metal filaments **30** in pores **28₁** overlying lower non-insulating region **22** as indicated in FIGS. **5h** and **6h**. The deposition is performed in the manner prescribed above in connection with the structure of FIGS. **1e** and **2e**. Since there is no electrical contact at the bottoms of pores **28₂** located directly above substrate **20**, substantially none of the filament metal accumulates in pores **28₂**. In FIG. **6h**, the dark circles indicate pores **28₁** filled with filaments **30**, while the light circles indicate empty pores **28₂**. Although there are openings **54** for all of pores **28**, the electrochemical nature of the process enables the deposition to be selective.

Filaments **30** again have the characteristics described above. Since pores **28₁** are situated at random locations across porous layer **24**, filaments **30** are again located in random places across layer **24**. Due to the centering that results from the track formation and etching, filaments **30** are self-aligned to openings **54₁** and therefore to gate electrode **46A**.

The cathode/gate structure of FIGS. **5h** and **6h** can be utilized directly as a field emitter. Nonetheless, it is again advantageous for filaments **30** to have sharpened tips that protrude out of the porous material. Accordingly, the structure of FIGS. **5h** and **6h** is further processed in the way described above for the other gated field emitters of the invention.

Portions of porous track layer **24A** exposed through gate openings **54₁** are removed with an isotropic etchant to form cavities **56₁** around filaments **30** as indicated in FIGS. **5i** and **6i**. Because gate layer **46A** also has openings **54₂**, portions of layer **24A** exposed through openings **54₂** are simultaneously removed to form cavities **56₂**. Cavities **56₁** and **56₂** (collectively “**56**”) can extend partway, or all the way, down to lower non-insulating region **22**. FIG. **5i** indicates the former case. In both cases, filaments **30** now protrude outward beyond the remainder **24F** of porous track layer **24A**.

The upper ends of filaments **30** are sharpened by performing an electropolishing and etching operation. An electropolishing step to round the edges of patterned gate layer **46A** overlying cavities **56₁** completes the basic fabrication of the field emitter. FIGS. **5j** and **6j** show the final structure

in which items **30B** are the sharpened remainders of filaments **30**. Item **46B** again is the edge-rounded remainder of gate layer **46A**.

Because electron-emissive elements **30B** are formed by a process that involves electrochemically filling pores **28₁** with metal and then (as desired) removing part of the metal to form sharpened tips, the upper ends of elements **30B** are normally situated below the bottom of patterned gate layer **46A** or **46B**. In certain applications, it is desirable for the electron-emissive elements in a gated field emitter to extend upward beyond the bottom of the gate electrode and sometimes even beyond the top of the gate electrode. Turning to FIGS. **7a–7e** (collectively “**FIG. 7**”), they illustrate a sequence of steps for converting filaments **30** in the structure of FIGS. **5i** and **6i** into pointed electron-emissive filaments that extend beyond the top of the gate electrode. FIG. **5i** is repeated here as FIG. **7a**.

A thin masking layer **86** of a lift-off material is formed on the partially finished field-emission structure of FIG. **7a** in such a way that the lift-off material fully covers gate layer **46A** but does not cover the tops of filaments **30**. See FIG. **7b**. Preferably, none of the lift-off material is present in the portions of cavities **56₁** situated to the sides of filaments **30**. The thickness of layer **86** is typically a fraction of (i.e., less than) the mean diameter of gate openings **54**.

Lift-off layer **86** is typically created according to a physical deposition technique, such as evaporation, in which atoms of the lift-off material impinge, and thereby accumulate, on the top of gate layer **46A** in a direction that is at a small angle to the (unshown) lower surface of substrate **20** and thus at a small angle to the top of gate layer **46A**. To achieve a spatially uniform deposition, either the partially finished field emitter or the source of the lift-off material is rotated at a constant speed about an axis perpendicular to the lower surface of substrate **20**. The impingement angle is set at a sufficiently small value as to enable some of the lift-off material to accumulate on the lateral edges of gate layer **46A** along gate openings **54**, but to substantially inhibit any of the lift-off material from passing through openings **54**.

Next, blunted metallic tips **88A₁** are respectively formed on the upper surfaces of filaments **30** as depicted in FIG. **7c**. Metal tips **88A₁** are preferably created according to a physical deposition technique, such as evaporation, in which atoms of the tip metal are directed toward the top of the field-emission structure in a direction largely perpendicular to the lower surface of substrate **20** and thus in a direction largely perpendicular to the top of gate layer **46A**. As a result, atoms of the tip metal pass through gate openings **54₁** to form tips **88A₁**.

During the tip deposition, annular portions **88B₁** of the tip metal respectively accumulate around filaments **30** at the bottoms of cavities **56₁**. Atoms of the tip metal also pass through gate openings **54₂** (not shown) to form pieces (not shown) of the tip metal along parts of insulating layer **24F** exposed through cavities **56₂** (also not shown). These pieces of the tip metal do not contact lower non-insulating region **22**. Accordingly, they do not affect the electrical operation of the final field emitter.

A layer **88C** of the tip metal accumulates on lift-off layer **86** during the tip deposition. As the thickness of layer **88C** increases, layer **88C** typically grows slightly in the lateral direction. Although this lateral growth gradually reduces the size of the openings through which the tip metal can deposit on filaments **30** and does affect the shape of metal portions **88B₁**, the deposition of the tip metal is terminated before the

tip-deposition openings through layer **88C** become so small that the tip metal starts to accumulate non-uniformly (e.g., to start forming conically-ended tips) on the upper surfaces of filaments **30**.

The combination of each filament **30** and overlying tip **88A₁** forms a composite lengthened electron-emissive filament. FIG. **8a** presents an enlarged view of one of lengthened filaments **30/88A₁** at the stage depicted in FIG. **7c**. The presence of metal portion **88B₁** along the length of filament **30/88A₁** is normally not electrically significant.

A noble metal is preferably employed for the tip metal. In a typical implementation, the tip metal consists of platinum when components **22**, **46A**, and **30** are respectively formed with chromium, molybdenum, and platinum. In this case, each filament **30** and overlying tip **88A₁** consist of the same metal. Alternatively, the tip metal can be a non-noble metal, such as molybdenum or niobium, capable of being physically deposited in the way described above. The amount of tip metal deposited depends on the desired length (or height) of the final electron-emissive elements.

Lift-off layer **86** is now removed by subjecting the structure to a chemical etchant that attacks the lift-off material but does not significantly attack any other materials in the structure. During the removal of layer **86**, layer **88C** is lifted off to produce the structure shown in FIG. **7d**.

An electropolishing and etching operation is performed in the manner described above to tailor and sharpen the upper ends of filaments **30/88A₁**. As in the fabrication process of FIGS. **1** and **2**, the electropolishing step enables filaments **30/88A₁** to become more uniform in length. During the filament etch, the upper ends of filaments **30/88A₁** are sharpened. FIG. **7e** illustrates the exemplary case in which substantially all the tailoring and sharpening is done on upper ends **88A₁** which thereby become pointed tips **88D₁**. Each cavity **56₁** in combination with underlying pore **28A₁** now forms a dielectric open space for corresponding filament **30/88D₁**. FIG. **8b** presents an enlarged view of typical filament **30/88D₁** at the stage shown in FIG. **7e**.

A further electropolishing step is normally also performed in the manner described above to round the edges of gate layer **46A** overlying cavities **56₁**. Patterned gate electrode **46B** in FIG. **7e** is the rounded-edge remainder of gate layer **46A**.

The mean diameter of the base of each metal tip **88D₁** in FIG. **7e** is the same as the mean diameter of underlying filament **30**. Alternatively, filamentary electron-emissive elements with conical tips can be created in which the mean diameter at the base of each tip is considerably greater than the mean diameter of the underlying filament. FIGS. **9a–9h** (collectively “FIG. **9**”) illustrate a sequence of fabrication steps by which the process of FIGS. **5** and **6** is modified starting at the stage shown in FIGS. **5f** and **6f** to create a gated field emitter having conically pointed filamentary electron-emissive elements whose tips are respectively wider at their bases than the underlying filamentary segments. FIG. **5f** is repeated here as FIG. **9a**.

Portions of insulating track layer **24** exposed through gate openings **54₁** are removed with an isotropic chemical etchant to form cavities **94₁** that extend partway, but not all the way, through layer **24** as shown in FIG. **9b**. Portions of layer **24** exposed through gate openings **54₂** (not shown) are simultaneously removed to form cavities **94₂** (likewise not shown) that extend partway through layer **24**. Each of cavities **94₁** and **94₂** (collectively “**94**”) extends slightly under gate layer **46A**.

The etchant preferably attacks the damaged material along track segments **50A** at approximately the same rate as

the undamaged insulating material of track layer **24**. Consequently, the portion of each track segment **50A**, of which only segments **50A₁** are shown in FIG. **9b**, at the bottom of corresponding cavity **94** is not significantly attacked during the etch. The remainder of insulating track layer **24** is indicated as item **24G** in FIG. **9b**.

The structure is now brought into contact with a chemical etchant that attacks the damaged **50A** track material much more than the undamaged material of insulating track layer **24G**. Pores **96₁** are thereby created through layer **24G** along the remaining portions of track segments **50A₁** at the bottom of cavities **94₁** as shown in FIG. **9c**. Pores **96₂** (not shown) are simultaneously etched through layer **24G** along the remaining (unshown) portions of track segments **50A₂** at the bottom of cavities **94₂** (also not shown). Item **24H** in FIG. **9c** is the remainder of insulating track layer **24G**.

The etch utilized to form pores **96₁** and **96₂** (collectively “**96**”) is performed uniformly in largely the same manner as that employed to create pores **28**. Each pore **96** is thereby centered on the location of corresponding track segment **50A**. Pores **96** typically have the same physical/spatial characteristics as given above for pores **28**. In addition to being narrower than overlying cavity **94**, each pore **96** is considerably narrower than corresponding gate opening **54**. Because (a) each track segment **50B** was in a straight line with corresponding track segment **50A** and (b) each gate opening **54** is centered on the location of corresponding track segment **50B**, each opening **54** is centered on corresponding pore **96**.

An appropriate filament metal is electrochemically deposited to create metal filaments **98** that substantially fill pores **96₁** over non-insulating region **22** as shown in FIG. **9d**. The filament deposition is performed in the manner used to create filaments **30** in the process of FIGS. **5** and **6**. Since there is no electrical contact at the bottoms of pores **96₂** located directly above substrate **20**, substantially none of the filament material accumulates in pores **96₂**. Consequently, the electrochemical nature of the process enables the deposition to be selective even through there is a gate opening for each pore **96**.

Filaments **98** can be formed with a noble metal such as platinum, gold, or palladium. The filament metal can also be a non-noble metal such as nickel, copper, cobalt, molybdenum, or niobium.

Metal filaments **98** have the same basic length, diameter, length-to-diameter, and cross-sectional characteristics as filaments **30**. In particular, the length-to-maximum-diameter ratio for filaments **98** is preferably 5 or more. Although a comparison of FIG. **9d** to FIG. **5i** might seem to indicate that filaments **98** are shorter than filaments **30**, filaments **98** can be made longer than filaments **30** by increasing the thickness of track layer **24** and appropriately controlling the depth of the cavity etch. As with filaments **30**, the centering that results from the track formation and etching enables filaments **98** to be self-aligned to gate openings **54₁** and thus to gate layer **46A**.

A thin masking layer **100** of a lift-off material is formed on the field-emission structure in such a manner that the lift-off material fully covers gate layer **46A** but does not cover the tops of filaments **98**. See FIG. **9e**. Lift-off layer **100** is preferably created according to the technique used to create lift-off layer **86** in the fabrication process of FIG. **7**.

A tip metal is introduced through gate openings **54₁** into cavities **94₁** to form pointed, generally conical metallic tips **102₁** that respectively contact filaments **98** as indicated in FIG. **9f**. The tip metal is normally the same as in the process of FIG. **7**.

The mean base diameter of each metallic tip 102_1 is normally greater than the diameter of underlying filament 98 . Consequently, tips 102_1 extend laterally over adjoining portions of track layer $24H$. The mean base diameter of each tip 102_1 , can be as large as, or slightly larger than, the mean diameter of corresponding gate opening 54_1 . Each filament 98 and overlying tip 102_1 form a composite electron-emissive element whose maximum diameter occurs at a point between—i.e., spaced apart from—the ends of that element.

Metal tips 102_1 are typically created according to a double-source physical deposition technique in which (a) atoms of a tip metal deposit on top of the structure in a direction largely perpendicular to the (unshown) lower surface of substrate 20 and thus in a direction largely perpendicular to the upper surface of gate layer $46A$ and (b) atoms of a closure material simultaneously impinge, and thereby accumulate, on the upper surface of layer $46A$ in a direction that is at a small angle to the lower surface of the substrate 20 while either the partially finished field emitter or the sources of the tip and closure materials are rotated at constant speed about an axis perpendicular to the lower surface of substrate 20 in order to obtain a spatially uniform deposition. Both parts of the deposition are usually done by evaporation.

The impingement angle for the closure material, which helps close the openings through which the tip metal enters cavities 94_1 is sufficiently small that substantially none of the closure material passes through gate openings 54_1 to accumulate on filaments 98 or on portions of track layer $24H$ exposed through cavities 94_1 . Only the tip metal passes through openings 54_1 . The tip metal also passes through gate openings 54_2 (not shown) to form pieces (not shown) of the tip metal along parts of layer $24H$ exposed through cavities 94_2 (also not shown). These pieces of the tip metal are electrically inconsequential since they do not contact lower non-insulating region 22 .

A composite layer 104 of the tip and closure materials accumulates on lift-off layer 100 during the tip deposition. Because the impingement angle for the closure material is less than 90° , composite layer 104 expands laterally as its thickness increases so as to progressively close the openings through which the tip metal can accumulate to form tips 102_1 . The deposition is conducted for a time sufficiently long that the tip-deposition openings through layer 104 close fully, and layer 104 becomes a continuous film. Tips 102_1 are thus formed in the shape of cones respectively centered on filaments 98 . Accordingly, tips 102_1 are self-aligned to gate openings 54_1 . Each filament 98 and overlying metal cone 102_1 form a composite electron-emissive element.

Lift-off layer 100 and composite layer 104 are removed in a conventional manner. FIG. $9g$ illustrates the resulting structure.

The final step in the fabrication process is to round the edges of gate layer $46A$ overlying cavities 94_1 as shown in FIG. $9h$. The edge rounding is preferably done by an electropolishing step in the way described above. Again, item $46B$ in FIG. $9h$ is the rounded-edge remainder of gate layer $46A$. Each cavity 94_1 and underlying pore 96_1 form a dielectric open space for corresponding electron-emissive element $98/102_1$. Since pores 96_1 are distributed randomly across track layer $24H$, electron-emissive elements $98/102_1$ are situated at random locations above non-insulating region 22 .

The metal portions that connect lower non-insulating region 22 to conical tips $88D_1$ and 102_1 in the structures

produced according to the fabrication processes of FIGS. 7 and 9 have been described as filaments. However, these metal portions could simply be metallic pedestals that do not necessarily have the elongated characteristics of filaments, such as the filamentary characteristics described above for filaments 30 in the process of FIGS. 5 and 6 . For example, the length (or height) of a pedestal could be less than its diameter. In manufacturing a gated field emitter according to the process of FIGS. 5 and 6 as modified in accordance with the steps of FIG. 7 or 9 , each reference to a filament can be broadened to a pedestal.

The process of FIGS. 5 and 6 can be modified to create a gated field-emission structure in which the electron-emissive elements consist solely of cones. That is, no filaments or pedestals lie between lower non-insulating region 22 and the cones. FIGS. $10a-10e$ (collectively "FIG. 10 ") and FIGS. $11a-11e$ (collectively "FIG. 11 ") illustrate a sequence of steps for creating such a cone-based gated field emitter from the structure depicted in FIGS. $5f$ and $6f$. These two figures are respectively repeated here as FIGS. $10a$ and $11a$.

Using gate layer $46A$ as an etch mask, dielectric open spaces 114_1 are etched through insulating track layer 24 down to lower non-insulating region 22 at the locations of gate openings 54_1 . See FIGS. $10b$ and $11b$. Dielectric open spaces 114_2 are similarly formed through layer 24 at the locations of gate openings 54_2 . Each of dielectric open spaces 114_1 and 114_2 (collectively " 114 ") extends slightly under gate layer $46A$. Item $24M$ in FIG. $10b$ is the remainder of track layer 24 .

A thin masking layer 116 of a lift-off material is created on the structure so as to fully cover gate layer $46A$ as shown in FIGS. $10c$ and $11c$. Lift-off layer 116 extends slightly over the edges of gate layer $46A$ along openings 54 . Layer 116 is preferably created according to the technique employed to form lift-off layer 86 in the fabrication process of FIG. 7 .

A cone metal is introduced through gate openings 54_1 into cavities 114_1 to form generally conical electron-emissive elements 118 , on non-insulating region 22 . See FIGS. $10d$ and $11d$. Electron-emissive cones 118_1 are preferably formed according to the double-source physical deposition technique used to create filament tips 102_1 in the process of FIG. 7 . The tip metal in the process of FIG. 7 is now the cone metal. Some of the cone metal also passes through gate openings 54_2 to form metal portions 118_2 in cavities 114_2 . Metal portions 118_2 do not contact lower non-insulating region 22 and therefore do not affect the electrical operation of the final field emitter.

During the cone deposition, a composite layer 120 of the cone and closure materials accumulates on lift-off layer 116 in the same manner that composite layer 104 is formed in the fabrication process of FIG. 9 . The cone-deposition openings through which the cone metal enters cavities 114 close in a substantially uniform manner. As a result, tips 118_1 are formed in the shape of electron-emissive cones self-aligned to gate openings 54_1 . The mean base diameter of each cone 118_1 is the same as, or slightly larger than, the mean diameter of corresponding gate opening 54_1 .

Layers 116 and 120 are removed in a conventional manner. FIGS. $10e$ and $11e$ depict the resulting structure. Because openings 54_1 are situated at random locations across gate layer $46A$, electron-emissive cones 118_1 are located in random places above non-insulating region 22 . An electropolishing step may be performed in the manner described above to round the edges of the remainder $46A$ of gate layer 46 .

Various modifications involving the lift-off layers can be made to the processes of FIGS. 7–11. Instead of creating a lift-off layer by an angled deposition on top of gate layer 46A after cavities 56, 94, or 114 have been formed in insulating track layer 24, the lift-off layer can be provided at an earlier stage in the process. For example, the lift-off layer can be deposited on gate layer 46 at the beginning of the process before the deposition of further layer 48 in FIG. 5b. With the lift-off layer now situated between layers 46 and 48, the lift-off layer undergoes the same operations that track layer 48 undergoes except that the lift-off layer is not removed when the etched remainder 48A of layer 48 is removed. In fact, the lift-off layer could even consist of layer 48A.

FIGS. 12a–12j (collectively “FIG. 12”) depict the exemplary case in which the process of FIG. 10 (and FIG. 11) is modified to include a lift-off layer 121 deposited on gate layer 46 before the deposition of track layer 48. Subject to the inclusion lift-off layer 121, FIGS. 12a–12j repeat FIGS. 5a–5e which portray the steps leading up to the structure of FIG. 10a. FIG. 12a identically repeats the starting structure of FIG. 5a.

As indicated in FIG. 12b, lift-off layer 121 is deposited on gate layer 46 after which further layer 48 is deposited on lift-off layer 121. The exposure of the structure to charged particles to form charged-particle tracks 50A and 50B respectively through insulating layer 24 and track layer 48 is shown in FIG. 12c. With lift-off layer 121 in place, FIG. 12d depicts how an etch is performed along tracks 50B to form apertures 52₁ through track layer 48. The remainder 48A of track layer 48 is then utilized as an etch mask to create corresponding openings through lift-off layer 121. See FIG. 12e. With the remainder 121A of lift-off layer 121 serving as an etch mask, FIG. 12f illustrates the etch of gate layer 46 to form gate openings 54₁.

Remaining track layer 46A is then removed as indicated in FIG. 12g. However, lift-off layer 121A remains in place. Dielectric open spaces 114₁ are then formed in insulating layer 24 as shown in FIG. 12h. Electron-emissive cones 118₁ are subsequently formed according to the above-described double-source physical deposition technique, with the result that composite layer 120 accumulates on lift-off layer 121A. Finally, lift-off layer 121A and overlying layer 120 are removed in a conventional manner to produce the final structure of FIG. 12j. This structure is substantially identical to the final structure of FIG. 10e.

FIGS. 13.1 and 13.2 illustrate the starting point for manufacturing implementations of the present field emitter in which lower non-insulating region 22 consists of an electrically conductive layer 22A situated under an electrically resistive layer 22B. As shown in FIG. 13.2, each of the lines that form region 22 consists of segments of both of layers 22A and 22B. Conductive layer 22A consists of one or more of the electrically conductive materials described above for layer 22. Resistive layer 22B is typically formed with cermet or lightly doped polycrystalline silicon.

FIGS. 14.1–14.4 respectively depict how the final structures of FIGS. 5j, 7e, 9h, and 10e appear when lower non-insulating region 22 consists of conductive layer 22A and resistive layer 22B. The lower ends of electron-emissive elements 30B, 30/88D₁, 98/102₁, or 118₁ contact resistive layer 22B. The resistance between each electron-emissive element and layer 22A is at least 10⁶ ohms, typically 10⁸ ohms or more.

In certain applications, it is desirable that each electron-emissive element contain an electrically resistive portion for

improving emission uniformity and limiting emitter currents to prevent short-circuit and arcing failures. The resistive portion, preferably contacts lower non-insulating region 22. The resistance for the resistive portion is normally at least 10⁶ ohms, preferably 10⁸ ohms or more. More specifically, the resistance for the resistive portion is at least 10¹⁰ ohms. The resistive portions typically consist of cermet or lightly doped polycrystalline silicon.

FIG. 15.1 illustrates an example in which each electron-emissive filament 30B in FIG. 1k, 3f, or 5j consists of a lower electrically resistive portion 30C and an upper electrically conductive electron-emissive pointed tip 30D. FIG. 15.2 illustrates an example in which each electron-emissive cone 118₁ in FIG. 10e consists of a lower electrically resistive portion 118A₁ and an upper electrically conductive electron-emissive portion 118B₁. To provide the electron-emissive elements with resistive portions 30C or 118A₁, an initial part of the emitter deposition is performed with an electrically resistive material rather than the material(s) specified above. For filaments 30 in FIGS. 1, 3, or 5, either all or a lower part of each filament 30 is formed with electrically resistive material. The same applies to filaments 98 in FIG. 9.

The electron-emissive cones in the field emitters of the invention have, for simplicity, been illustrated in the drawings as being of right circular shape (i.e., a right triangle rotated about one of its legs). However, the cones invariably have surfaces that differ somewhat from a right circular shape. For example, the surfaces of the cones typically bend inward slightly as shown in FIG. 16.1 for cones 102₁ of FIG. 9h and as depicted in FIG. 16.2 for cones 118₁ of FIG. 10e.

FIGS. 17.1–17.4 illustrate several longitudinal shapes that filaments 30B can have in the final field emitters of FIG. 1k, 3f, and 5j. As indicated in FIGS. 17.1 and 17.2, filaments 30B can be solid cylinders except at their upper ends. If tracks 50 are etched in such a manner as to create pores 28 as inverted cones, filaments 30B can be inverted solid cones which taper down in transverse cross section from just below their upper ends to their lower ends in the manner shown in FIGS. 17.3 and 17.4. Depending on how the electropolishing and etching operation is done, the electron-emissive tips at the upper ends of filaments 30B can be generally rounded as indicated in FIGS. 17.1 and 17.3 or sharply pointed as indicated in FIGS. 17.2 and 17.4.

FIG. 18 depicts the electrochemical deposition system employed during the electrochemical filament deposition operations described above. The electrochemical system consists of an electrochemical cell 72 and a power supply and control system 74. In turn, electrochemical cell 72 consists of cell electrolyte 76, a surrounding sidewall 78, an O-ring 80, an anode 82, and a cathode 84 formed by part of the structure on which metal is being deposited. Cathode 84 includes lower non-insulating region 22 to which power supply and control system 74 is connected. FIG. 18 specifically illustrates the filament deposition in the process of FIGS. 1, 3, and 5.

The electrochemical system shown in FIG. 18 operates according to a constant-current technique. Alternatively, a constant-voltage electrochemical deposition system could be used.

In the field emitters described above, each gate opening exposes only one electron-emissive element. Alternatively, charged-particle tracks can be utilized in accordance with the invention to fabricate gated area field emitters in which the gate electrode has gate openings respectively centered on the charged-particle tracks and in which each gate opening

exposes multiple electron-emissive elements generally centered as a group on the gate opening.

FIGS. 19a–19d (collectively “FIG. 19”) and FIGS. 20a–20d (collectively “FIG. 20”) illustrate a sequence of additional processing steps that can be applied to the intermediate structure of FIGS. 5f and 6f to produce such a gated area field emitter in which each gate opening 54₁ accommodates a group of electron-emissive elements. FIGS. 5f and 6f are respectively repeated here as FIGS. 19a and 20a.

Inasmuch as each gate opening 54₁ exposes multiple electron-emissive elements in the process of FIGS. 19 and 20, gate openings 54₁ are usually wider here than in the earlier fabrication processes described above. In particular, the average mean diameter of openings 54₁ in FIGS. 19a and 20a is 0.5–5 μm, typically 1 μm. In comparison to using a photolithographic etching technique to form openings 54₁, the use of the present charged-particle track technique is particularly advantageous when the mean opening diameter is 1 μm or less. The average density of openings 54₁ is 10⁶–10⁸ openings/cm², typically 10⁷ openings/cm².

Using non-insulating layer 46A as a mask, insulating track layer 24 is etched through gate openings 54₁ to create corresponding dielectric open spaces 128₁ down to lower non-insulating region 22 as shown in FIGS. 19b and 20b. If gate openings 54₂ are present in layer 46A, the etchant also attacks portions of layer 24 exposed through openings 54₂ to create corresponding dielectric open spaces 128₂, indicated only in FIG. 20b, down to, and possibly partly into, insulating substrate 20. The etch is normally performed in such a way that dielectric open spaces 128₁ and 128₂ (collectively “128”) extend laterally under layer 46A in a generally uniform manner as indicated in FIG. 19b. However, the etch could be performed so that the sidewalls of open spaces 128 respectively line up with the edges of gate openings 54. In either case, each dielectric open space 128 is centered on overlying gate opening 54. Item 24P in FIGS. 19b and 20b is the remainder of track layer 24.

A group of preformed particles 130 are introduced into each dielectric open space 128₁ and then distributed in a relatively uniform manner across the portion of the upper surface of lower non-insulating region 22 at the bottom of open space 128₁. See FIGS. 19c and 20c. The distributing step is performed in such a way that particles 130 are laterally separated from one another and are securely fixed to, and electrically coupled to, non-insulating region 22.

The distributing step typically involves dispersing particles 130 across the bottom of open spaces 128₁ in a random manner and then performing an operation to bond particles 130 to non-insulating region 22. Electrically non-insulating particle bonding material (not shown) holds particles 130 in place. To the extent that particles 130 do not actually touch region 22, the bonding material electrically connects particles 130 to region 22. The average mean diameter of particles 130 is 5 nm–1 μm, typically 100 nm.

Particles 130 preferably are electron emissive prior to being introduced into dielectric open spaces 128₁. However, particles 130 could be introduced into open spaces 128₁ in a non-emissive state and then be made electron emissive during or after the bonding operation. In either case, particles 130 serve as electron-emissive elements.

In one embodiment, the dispersion of particles 130 across the exposed upper surface portions of non-insulating region 22 and the subsequent bonding of particles 130 to region 22 is performed in the manner described in Twichell et al, co-filed U.S. patent application Ser. No. 08/267,283, “Structure and Fabrication of Electron-emitting Devices Utilizing

Electron-emissive Particles which Typically Contain Carbon,” filed 29 Jun. 1984. Particles 130 then typically consist of at least 50 atomic percent carbon in the form of electrically non-insulating diamond, graphite, amorphous carbon, or/and electrically non-insulating silicon carbide.

During the introduction of particles 130 into open spaces 128₁, some preformed particles (not shown) may be introduced into dielectric open spaces 128₂. Should this happen, there will be no significant deleterious effect on device operation because any electron-emissive particles at the bottoms of openings 128₂ contact insulating material of substrate 20.

An electropolishing step can be performed to round the edges of patterned gate layer 46A. The final gated field-emitting device is shown in FIGS. 19d and 20d. Item 46B again indicates the edge-rounded remainder of gate layer 46A. Since dielectric open spaces 128₁ are respectively centered on gate openings 54₁, electron-emissive particles 130 in each open space 128₁ are, as a group, centered on overlying gate opening 54₁.

Particles 130 could be formed with materials other than carbon. For example, molybdenum or/and doped silicon carbide could be used. Instead of being dispersed across non-insulating region 22 in a preformed state, particles 130 could be grown over region 22.

FIGS. 21a–21e (collectively “FIG. 21”) illustrate another processing sequence that can be applied to the intermediate structure of FIGS. 5f and 6f to produce a gated field emitter in which each gate opening 54₁ accommodates multiple electron-emissive elements. FIG. 5f is repeated here as FIG. 21a except that non-insulating region 22 is thicker. In particular, the thickness of region 22 in the structure of FIG. 21a is 0.1–2 μm, typically 0.5 μm, greater than in the structure of FIG. 5f. The average mean diameter of gate openings 54₁ here is 0.5–5 μm, typically 1 μm. The average density of openings 54₁ again is 10⁶–10⁸ openings/cm², typically 10⁷ openings/cm².

Insulating track layer 24 is etched through gate openings 54₁ to create dielectric open spaces 128₁ down to lower non-insulating region 22 as depicted in FIG. 21b. A group of preformed particles 130 are introduced into each dielectric open space 128₁ and then distributed uniformly across region 22 at the bottoms of open spaces 128₁ so that particles 130 again are laterally separated from one another and are securely fixed, and electrically coupled to, region 22. See FIG. 21c. The distributing step is performed in the manner described above. If particles 130 are not electron emissive prior to being introduced into open spaces 128₁, particles 130 can be made electron-emissive during or after bonding to region 22.

Material of non-insulating region 22 not covered (or not shadowed) by particles 130 is removed to a depth less than the thickness of region 22 to create electrically non-insulating pedestals 132 respectively below particles 130 as shown in FIG. 21d. The removal operation is preferably done in the manner described in Twichell et al, cited above. In particular, the removal operation typically entails milling or anisotropically etching region 22 using particles 130 as masks. The side surface of each pedestal 132 extends vertically or, in going from top to bottom (i.e., downward), slopes inward to a point at or just above the bottom of pedestal 132.

The mean height of pedestal 132 is 0.1–2 μm, typically 0.5 μm. The combination of each pedestal 132 and overlying electron-emissive particle 130 constitutes an electron-emissive pillar. Item 22C in FIG. 21d is the remainder of lower non-insulating region 22.

An electropolishing step can again be performed to round the edges of gate layer 46A. FIG. 21e shows the final gated field emitter. Because dielectric open spaces 128₁ are centered on gate openings 54₁, electron-emissive pillars 130/132 in each open space 128₁ are, as a group, centered on overlying gate opening 54₁.

FIGS. 22a–22c (collectively “FIG. 22”) illustrate part of a procedure for manufacturing a gated field emitter having a structure very similar to that of FIGS. 19d and 20d. FIG. 22a depicts a fabrication stage analogous to the stage shown in FIG. 19a, and thus the stage shown in FIG. 5f, except that a layer of laterally separated particles 134 is situated along the interface between lower non-insulating region 22 and insulating track layer 24. The structure of FIG. 22a is created according to the fabrication process of FIGS. 5 and 6 except that preformed particles 134 are distributed across the upper surface of region 22 prior to the deposition of insulating track layer 24—i.e., between the stages shown in FIGS. 5a and 5b.

Particles 134 are distributed across non-insulating region 22 in such a way that they are securely fixed to, and electrically coupled to, region 22. The distribution step typically entails dispersing particles 134 across region 22 in a random manner and then bonding particles 134 to region 22 using electrically non-insulating particle bonding material (not shown). Particle 134 preferably have the characteristics described above for particles 130. In one embodiment, the distribution of particles 134 across region 22 is performed as disclosed in Twichell et al, cited above.

Using non-insulating layer 46A as a mask, insulating track layer 24 is etched through gate openings 54₁ substantially down to lower non-insulating region 22 to form corresponding dielectric open spaces 136₁ as shown in FIG. 22b. A group 134A of particles 134 is thereby exposed at the bottom of each open space 136₁. If gate openings 54₂ are present in non-insulating layer 46A, track layer 24 is simultaneously etched through gate openings 54₂ to create electrically inconsequential dielectric open spaces 136₂ (not shown) down to, and possibly partially into, insulating substrate 20. Dielectric open spaces 136₁ and 136₂ (collectively 136) normally extend laterally under gate layer 46A generally in the illustrated manner but could have their sidewalls vertically aligned with the edges of layer 46A.

Particles 134 preferably are electron emissive before being dispersed across lower non-insulating region 22. Alternatively, at least particles 134A can be made electron emissive at some later point, including subsequent to the formation of dielectric open spaces 136. In either case, particles 134A serve as electron-emissive elements. The edges of patterned gate layer 46A can be rounded by an electropolishing operation. The final gated field emitter shown in FIG. 22c is quite similar to that of FIG. 19a.

The variations described above with respect to particles 130 can generally be applied to particles 134. Also, subject to increasing the thickness of lower non-insulating region 22, the structure of FIG. 22c can be further processed according to the steps described for FIGS. 21d and 21e to produce a gated field emitter having pillared electron-emissive elements consisting of particles 134 on underlying pedestals.

FIGS. 23a–23h (collectively “FIG. 23”) and FIGS. 24a–24d (collectively “FIG. 24”) illustrate a process for manufacturing a gated area field-emission cathode structure using charged-particle tracks and emitter etching according to the teachings of the invention. The field-emission structure of FIGS. 23 and 24 is typically utilized to excite

phosphors on a faceplate in a CRT of a flat-panel display such as a flat-panel television screen or video monitor.

The starting point for the fabrication process is a substrate 140 typically created from a plate having a largely flat upper surface and a largely flat lower surface (not shown) extending substantially parallel to the upper surface. See FIG. 23a. Substrate 140 normally consists, at least along its upper surface, of electrically resistive (intrinsic or lightly doped) semiconductor material or/and electrically insulating material. The resistive semiconductor material preferably is silicon but can be germanium or gallium arsenide. The insulating material is ceramic or/and glass.

An electrically non-insulating emitter layer 142 is provided along the top of substrate 140 as indicated in FIG. 23a. Emitter layer 142 preferably consists of an electrical conductor, specifically conductively doped semiconductor material or/and metal. The conductively doped semiconductor material typically is silicon of n-type or p-type conductivity but can be germanium or gallium arsenide. When substantially all of layer 142 is conductively doped silicon, the emitter thickness is 0.1–1 μm, typically 0.2 μm. For the case in which metal is used to form layer 142, the metal is typically titanium.

Emitter layer 142 is configured and used in largely the same way as lower non-insulating region 22 above. In particular, layer 132 is typically a patterned layer containing a group of parallel emitter lines.

If substrate 140 consists of electrically resistive semiconductor material at least along the upper substrate surface, the emitter lines typically consist of conductively doped semiconductor material created by selectively introducing suitable dopant into the resistive semiconductor material. FIG. 25a depicts a vertical cross section through the structure of FIG. 23a for such an embodiment. FIG. 26a depicts a vertical cross section through the structure of FIG. 23a for an embodiment in which the emitter lines consist of metal or conductively doped semiconductor material formed on top of substrate 140.

An electrically insulating track (or track-recording) layer 144 is formed on top of the structure. Parts of track layer 144 are situated on both substrate 140 and emitter layer 142. The thickness of layer 144 is 0.1–2 μm, typically 0.5 μm, depending on the desired diameter of apertures later formed through layer 144. Suitable materials for layer 134 are the same as those listed above for insulating track layer 24.

The structure is subjected to energetic charged particles that impinge on top of track layer 144 in a direction largely perpendicular to the (unshown) flat lower surface of substrate 140 and thus in a direction generally perpendicular to the upper structural surface. The charged particles have sufficient energy to pass fully through layer 144 so as to form straight tracks through layer 144 at random locations across layer 144. FIGS. 23b and 24a illustrate the track formation. The charged-particle tracks constitute damaged zones along the particle paths.

The charged-particle tracks are indicated by reference symbols beginning with “146” in FIGS. 23b and 24a. Although the charged particles also pass through emitter layer 142 (and typically into substrate 140), the charged particles do not significantly damage layer 142 and therefore do not create charged-particle tracks through layer 142. Two of the lines that typically form layer 142 are shown in dashed form in FIG. 24a. As indicated there, the tracks fall into two categories: (a) tracks 146₁ extending through portions of layer 144 overlying emitter layer 142 and (b) tracks 146₂ extending through portions of layer 144 situated directly on substrate 140 to the sides of layer 142.

As with charged-particle tracks **26** above, charged-particle tracks **146**₁ and **146**₂ (collectively “**146**”) extend parallel to one another in a direction generally perpendicular to the upper structural surface. Tracks **146** have similar characteristics, to, and are formed according to the same techniques as, tracks **26**. The density of tracks **146** likewise is 10^6 – 10^9 tracks/cm², typically 10^7 – 10^8 tracks/cm². For illustrative purposes, only a small portion of tracks **146** is indicated in FIGS. **23b** and **24a**.

The damaged insulating material along tracks **146** is removed by bringing track layer **144** into contact with a suitable chemical etchant that attacks the damaged track material much more than the undamaged material of layer **144**. As a result, generally circular pores are etched through layer **144** along tracks **146** down to emitter layer **142**. The etchant preferably does not significantly attack any other parts of the field-emission structure.

The etch is continued into the largely undamaged material of track layer **144** to broaden the pores. Apertures **148**₁ and **148**₂ (collectively “**148**”) are thereby respectively created along tracks **146**₁ and **146**₂. See FIGS. **23c** and **24b**. Apertures **148**₁ expose corresponding portions of the upper surface of emitter layer **142**.

The full etch of track layer **144** is performed in a laterally uniform manner. Accordingly, each aperture **148** is generally circular in plan view as indicated in FIG. **24b**. Each aperture **148** is also centered on corresponding track **146**. The thickness of layer **144** is reduced during the etch.

The second part of the insulating-material etch can be done with the etchant used during the first part or with another etchant. In either case, components **140** and **142** are not significantly attacked during the second part of the etch. When track layer **144** consists of a polymer, both parts of the etch are preferably done with sodium hydroxide or potassium hydroxide.

Apertures **148** reach an average mean diameter of 0.1–2 μm , typically 0.2 μm , along the bottom of the reduced-thickness remainder **144A** of track layer **144**. The aperture diameter is substantially the same for all of apertures **148**. For illustrative purposes, the lateral dimensions of apertures **148** compared to the widths of the lines that form emitter layer **142** are greatly exaggerated in the plan-view drawings.

Apertures **148**₂, which extend through portions of track layer **144A** lying directly above substrate **140**, do not significantly affect device operation. Accordingly, apertures **148**₂ are not discussed further below or shown in any of the remaining drawings. In fact, the creation of apertures **148**₂ could be avoided by using an appropriate mask during either the track-formation step or the aperture-etch step.

A cap layer **150** is deposited on top of the structure as shown in FIG. **23d**. Cap layer **150** contains (a) main cap regions **150A** situated on emitter layer **142** at the bottoms of apertures **148**₁ and (b) a further cap region **150B** situated on top of track layer **144A**. Cap layer **150** may be formed with electrically insulating material or with electrically non-insulating material (or even with both types of material). For example, layer **150** typically consists of a metal such as chromium when emitter layer **132** is conductively doped silicon.

Cap layer **150** may be deposited by evaporating or sputtering the desired cap material. Alternatively, main cap regions **150A** can be formed by a selective deposition technique such as electrochemical deposition. In this case, substantially none of the cap material accumulates on track layer **144A**—i.e., further cap region **150B** is not formed.

The thickness of cap layer **150** is less than the thickness of track layer **144A**. Specifically, the cap thickness is 0.05–1

μm , typically 0.2 μm . Although not shown in FIG. **23d**, small pieces of the cap material may accumulate along the sidewalls of track layer **144A** above main cap regions **150A**. To the extent that such sidewall cap pieces are formed and not removed during the track-material dissolving operation described below, these sidewall cap pieces are removed according to a conventional technique that may slightly reduce the thickness of regions **150A** and **150B**.

The structure is subjected to an agent that dissolves track layer **144A** but does not significantly affect cap layer **150** or any of the other structural components. All of layer **144A** is removed during the etch. Further cap region **150B** (when present) is lifted off during the removal of layer **144A** to produce the structure shown in FIG. **23e**. When layer **144A** consists of a polymer such as polycarbonate, the dissolving step is performed with chloroform. A dilute hydrofluoric acid solution is used as the dissolving agent when layer **144A** is glass. Since remaining cap regions **150A** were formed in apertures **148**₁, cap regions **150A** are respectively centered on the locations of charged-particle tracks **146**₁.

Next, the structure is etched with an etchant that attacks emitter layer **142** but does not significantly attack cap regions **150A**. The emitter etch is performed in such a way as to uniformly remove (a) emitter material not covered by cap regions **150A** and (b) laterally adjacent emitter material extending partway under regions **150A**, thereby creating a depression **152** in layer **142**. See FIGS. **23f** and **24c**. Regions **150A** act as etch masks to control the lateral extent of the etch. The etchant preferably is a reactive-ion etchant.

The emitter etch is conducted for a time sufficiently long to underetch a large fraction of lower surface of each cap region **150A** but not long enough for depression **152** to reach substrate **140** or for the tops of the sidewalls of depression **150** to form points below regions **150A**. As a result, regions **150A** remain in place. Item **142A** in FIG. **23f** is the remainder of emitter layer **142**. Along the upper surface of remaining emitter layer **142A**, depression **152** defines truncated generally conical emitter portions **142B** in layer **142A**. Because the etch is done uniformly, each emitter portion **142B** is centered on, and thereby aligned to, overlying cap region **150A**.

Emitter portions **142B** are sharpened by reacting emitter material along the upper surface of layer **142A** with one or more other materials to form a layer **154** consisting of a compound of these materials. FIG. **23g** shows the resultant structure. Item **142C** is the remainder of emitter layer **142A**. The reaction process consumes an amount of emitter material sufficient to enable generally conical electron-emissive portions **142D** to be defined in the emitter material along the upper surface of layer **142C**. Each emitter portion **142D** has a sharply pointed tip directed towards a corresponding one of cap regions **150A**.

Emitter portions **142D** are generally cones even though their side surfaces are concave as viewed from the side. Each electron-emissive cone **142D** is centered on overlying cap region **150A** and therefore on the location of corresponding charged-particle track **146**₁. Cones **142D** have an average height of 0.1–2 μm , typically 0.2 μm . The base diameter of cones **142** is approximately the same as that of cap regions **150A** and, accordingly, approximately the same as that of apertures **148**₁. That is, cones **142D** have an average mean base diameter of 0.1–2 μm , typically 0.2 μm .

Compound layer **154** is preferably an oxide of the emitter material. The emitter oxide is typically created by exposing the structure to an oxygen-containing gas, such as wet or dry oxygen, at high temperature. When emitter layer **142C** is

conductively doped silicon, layer **154** is silicon oxide. Likewise, layer **154** is a metal oxide when layer **142C** is metal.

Electrically insulating material is deposited on the structure by causing the constituents of the insulating material to move towards the upper surface of the structure in a direction largely perpendicular to the lower structural surface. As shown in FIG. **23h**, portions **156A** of the insulating material accumulate on cap regions **150A**. A portion **156B** of the insulating material accumulates on the portion of compound layer **154** not covered (or not shadowed) by regions **150A**. Insulating portions **156A** and **156B** together constitute a discontinuous insulating layer **156**. The deposition of insulating layer **156** is typically performed by sputtering or chemical vapor deposition. Layer **156** typically consists of silicon oxide.

Next, electrically non-insulating gate material is similarly deposited on the upper surface of the structure in a direction generally perpendicular to the lower structural surface. As also shown in FIG. **23h**, portions **158A** and **158B** of the gate material respectively accumulate on insulating portions **156A** and **156B**. Gate portions **158A** and **158B** together form a discontinuous gate layer **158**. The deposition of gate layer **158** is typically performed by sputtering or evaporation. Layer **158** usually consists of an electrical conductor, typically a metal such as molybdenum. Layer **158** could also be formed with conductively doped semiconductor material such as n-type or p-type polycrystalline silicon. The composite thickness of insulating layer **156** and gate layer **158** is less than the height of emitter cones **142D**.

Cap regions **150A** are removed by subjecting the structure to an etchant that attacks the exposed material of compound layer **154** under cap regions **150A** but does not significantly attack the gate material or emitter layer **142C**. For example, a buffered hydrofluoric acid solution can again be employed when layer **154** consists of silicon oxide. The portions of layer **154** lying below cap regions **150A** are thereby removed. As a result, regions **150A** are lifted off along with overlying insulating portions **156A** and gate portions **158A**. The etching of layer **154** normally extends slightly under gate portions **156A**. FIG. **23i** depicts the resultant structure in which item **154A** is the remainder of layer **154**.

The etchant typically attacks the side edges of insulating portions **156B** so as to slightly undercut remaining gate portion **158B**. Item **156C** in FIG. **23i** is the remainder of portion **156B**. Of course, layer **156B** will remain fully in place to support gate portion **158B** if the etchant does not attack layer **156B**.

Remaining gate portion **158B** forms a patterned gate layer. Emitter cones **142D** preferably extend partially through gate openings **160** in patterned gate layer **158B**. Because cap regions **150A** were respectively centered on cones **142D**, gate openings **160** are respectively centered on, and thus aligned to, cones **142D**.

Remaining insulating portion **156C** (or **156B**) forms a patterned insulating layer through which dielectric openings **162** extend. If remaining compound portion **154A** consists of insulating material, portion **154A** forms part of this patterned insulating layer. As with gate openings **160**, dielectric openings **162** are centered on cones **142D** as the result of the self alignment that occurs during the fabrication process. Since each cone **142D** is centered on the location of corresponding charged particle track **146₁**, each gate opening **160** and underlying dielectric opening **162** are centered on the location of corresponding track **146₁**.

Using a suitable photoresist mask (not shown) gate layer **158B** is patterned into a group of lines extending perpen-

dicular to the lines that form emitter layer **142C**. FIGS. **23j** and **24d** depict the final structure in which item **158C** is the patterned remainder of gate layer **158D**. FIG. **24d** illustrates one of the lines that form patterned gate layer **158C**. Emitter cones **142D** are electron-emissive elements which, in combination with the underlying structural components, form a gated field emitter.

FIGS. **25b** and **26b** illustrate typical vertical cross sections through the final structure of FIGS. **23j** and **24d**. FIG. **25b** represents the embodiment where emitter lines **142** are conductively doped regions created in electrically resistive semiconductor material. FIG. **26b** represents the embodiment where lines **142** consist of metal or conductively doped semiconductor material formed on substrate **140**.

As with lower non-insulating region **22** in the earlier-described field emitters of the invention, emitter layer **142** in the field emitter of FIGS. **23j** and **24d**, could be provided as a lower electrically conductive sublayer and an upper electrically resistive sublayer. The conductive sublayer would be formed with one or more of the electrical conductors described above for layer **142**. The resistive sublayer would typically consist of cermet or lightly doped polycrystalline silicon.

The field emitters of the present invention operate in the following way when used in a flat-panel CRT display where emitter layer **142** (in the embodiment of FIGS. **23j** and **24d**) or lower non-insulating region **22** (in the other embodiments) contains emitter lines that are situated below, and cross, lines of the gate layer. An anode (or collector) structure is situated a short distance away from the top of each emitter. The anode is maintained at high positive voltage relative to the gate and emitter lines.

When a suitable voltage is applied between (a) a selected one of the gate lines and (b) a selected one of the emitter lines, the selected gate line extracts electrons from the electron-emissive elements at the intersection of the two selected lines and controls the magnitude of the resulting electron current. Desired levels of electron emission typically occur when the applied gate-to-emitter electric field reaches 20 volt/ μm or less at a current density of 1 mA/cm² as measured at the phosphor-coated faceplate of the flat-panel display. The extracted electrons are subsequently collected at the anode.

Directional terms such as "lower" and "down" have been employed in describing the present invention to establish a frame of reference by which the reader can more easily understand how the various parts of the invention fit together. In actual practice, the components of a field emitter may be situated at orientations different from that implied by the directional terms used here. The same applies to the way in which the fabrication steps are performed in the invention. Inasmuch as directional terms are used for convenience to facilitate the description, the invention encompasses implementations in which the orientations differ from those strictly covered by the directional terms employed here.

The various electron-emissive elements and charged-particle tracks (or track segments) have longitudinal axes (not shown). Each electron-emissive element is generally symmetric about its longitudinal axis. A reference to an etch as being performed along a charged-particle track through a track layer means that the removed material occupied a volume containing at least part of the track's longitudinal axis in the track layer.

While the invention has been described with reference to particular embodiments, this description is solely for the purpose of illustration and is not to be construed as limiting

the scope of the invention claimed below. For example, the gated area electron emitters of the invention have been described as operating according to the field-emission mode in which selected parts of the patterned gate electrode are furnished with voltage sufficient to extract electrons from selected electron-emissive elements. Nonetheless, another mechanism such as photoemission or thermionic emission could be employed to cause part or all of the electron emission. In such embodiments, selected parts of the gate electrode typically collect electrons to cut off electron flow to corresponding parts of the anode.

Under certain conditions, the double-source deposition utilized to form conical tips **102₁** in the fabrication process of FIG. **9** can be replaced with a single-source physical vapor deposition in which only the tip material is deposited. No separate closure material is employed. The same applies to the double-source deposition used to form cones **118₁** in the process of FIGS. **10** and **11**.

Instead of being formed with metal, tips **88D₁**, and cones **102₁** and **118₁** could be formed with other electrical conductors such as conductively doped semiconductors. Components **88D₁**, **102₁**, and **118₁** along with gate layers **34**, **40**, **46**, and **158** could be formed with electrically resistive material such as lightly doped semiconductors. Gate layer **34**, **40**, or **46** could be patterned into lines running perpendicular to the lines that form region **22**.

Each of gate electrodes **34B**, **40B**, **46B** (or **46A**), and **158C** could be employed to modulate the movement of electrons extracted by the anode. Various modifications and applications may thus be made by those skilled in the art without departing from the true scope and spirit of the invention as defined in the appended claims.

We claim:

1. A method comprising the steps of:

creating a structure in which an electrically insulating layer lies over a lower electrically non-insulating region, an electrically non-insulating gate layer lies over the insulating layer, a lift-off layer lies over the gate layer, and a further layer lies over the lift-off layer; providing a multiplicity of apertures through the further layer;

etching the lift-off layer through the apertures to form corresponding lift-off openings through the lift-off layer;

etching the gate layer through the lift-off openings to form corresponding gate openings through the gate layer;

etching the insulating layer through the gate openings to form corresponding dielectric open spaces through the insulating layer;

forming a like multiplicity of electron-emissive elements over the lower non-insulating region such that each electron-emissive element contacts the lower non-insulating region through a corresponding one of the dielectric open spaces, electrically non-insulating material being deposited over the lift-off layer and into the dielectric open spaces so as to form at least part of each electron-emissive element; and

removing the lift-off layer so as to substantially remove any of the non-insulating material accumulated over the lift-off layer.

2. A method as in claim **1** wherein the providing step comprises:

causing charged particles to pass through the further layer to form a like multiplicity of charged-particle tracks therethrough; and

etching the further layer along the tracks to form the apertures.

3. A method as in claim **1** wherein the second-mentioned etching step includes etching the gate layer through the apertures in the further layer.

4. A method as in claim **1** further including, prior to the removing step, the step of removing the further layer.

5. A method as in claim **1** wherein the further layer consists primarily of at least one of (a) organic polymer, (b) glass, and (c) crystalline material.

6. A method as in claim **1** wherein each electron-emissive element comprises electrically non-insulating material generally in the shape of an electron-emissive cone that points away from the lower non-insulating region.

7. A method as in claim **1** wherein each electron-emissive element further includes an electrically non-insulating pedestal situated between the lower non-insulating region and that element's cone.

8. A method as in claim **1** wherein each electron-emissive element is generally in the shape of a filament.

9. A method as in claim **1** wherein each electron-emissive element comprises (a) an electrically resistive portion situated over the lower non-insulating region and (b) an electrically conductive portion situated over the resistive portion.

10. A method as in claim **1** wherein the lower non-insulating region comprises an electrically conductive part and an electrically resistive part situated over the conductive part.

11. A method as in claim **1** wherein each lift-off opening is generally centered on the corresponding aperture, each gate opening is generally centered on the corresponding lift-off opening, and each dielectric open space is generally centered on the corresponding gate opening.

12. A method as in claim **1** wherein the electron-emissive elements are operable in field-emission mode.

13. A method as in claim **1** wherein the forming step entails depositing the non-insulating material for a time sufficient to allow the non-insulating material to accumulate in the dielectric open spaces generally in the shape of cones that respectively form the electron-emissive elements.

14. A method as in claim **13** wherein the further layer consists primarily of polycarbonate.

15. A method comprising the steps of: providing a multiplicity of apertures through a track layer; etching an underlying lift-off layer through the apertures to form corresponding lift-off openings through the lift-off layer;

etching an underlying electrically non-insulating gate layer through the lift-off openings to form corresponding gate openings through the gate layer;

etching an underlying electrically insulating layer through the gate openings to form corresponding dielectric open spaces through the insulating layer down to an underlying lower electrically non-insulating region;

depositing electrically non-insulating material over the lift-off layer and into the dielectric open spaces such that the non-insulating material accumulates in the dielectric open spaces generally in the shape of corresponding electron-emissive elements that contact the lower non-insulating region; and

subsequently removing the lift-off layer so as to substantially remove any material overlying the lift-off layer.

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16. A method as in claim **15** wherein the providing step comprises:
causing charged particles to pass through the track layer to form a like multiplicity of charged-particle tracks therethrough; and
etching the track layer along the tracks to form the apertures.

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17. A method as in claim **16** wherein the track layer consists primarily of at least one of (a) organic polymer, (b) glass, and (c) crystalline material.

18. A method as in claim **16** wherein the track layer⁵ consists primarily of polycarbonate.

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